A METHOD FOR DETERMINING GENETIC AFFILIATION, SUBSTRUCTURE AND GENE FLOW WITHIN HUMAN POPULATIONS

CROSS-REFERENCE

[0001] This application claims the benefit of U.S. Provisional Application No. 06/245,355, filed November 1, 2000, which application is incorporated herein by reference.

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH

[0002] This invention was made with government support under grant nos. GM55273 and GM 28428 awarded by the NIH. The government may have certain rights in this invention.

FIELD OF THE INVENTION

[0003] The present invention relates to nucleic acid polymorphisms and their methods of use in, for example, determination of paternity and forensics.

BACKGROUND OF THE INVENTION

[0004] The science of genetics has taken a keen interest in the identification of human individuals and genetic relationships between individuals. The genome of an individual is unique to that individual, and can be used for identification purposes, *e.g.,* testing for paternity and/or forensic testing (e.g. to identify an individual in the context of post-mortem identification or in the criminal justice system). Procedures have been developed which are based on identification and characterization of changes in an individual's DNA, referred to as DNA polymorphisms, where such changes are due to nucleotide substitution, insertion, or deletion within the chains of DNAs.

[0005] In forensics, for example, there is an interest in polymorphisms for identification purposes. Techniques have been developed to compare homologous segments of DNA to determine if the segments are identical or if they differ in one or more nucleotides. Practical applications of these techniques relate to fields other than forensic medicine, for example, genetic disease diagnosis and human genome mapping.

[0006] The most accurate and informative way to compare DNA segments requires a method which provides the complete nucleotide sequence for each DNA segment. Particular techniques have been developed for determining actual sequences in order to study mutation in human genes. See, for example, Proc. Natl. Acad. Sci. U.S.A. 85, 544-548 (1988) and Nature 330, 384-386 (1987). However, because of the extensive amounts of time and high costs to determine, interpret, and compare sequence information, presently it is not practical to use extensive sequencing for compare more than just a few DNA segments.

[0007] A frequently used technique for screening for DNA polymorphisms arising from mutations consist of digesting the DNA strand with restriction endonucleases and analyzing the resulting fragments by means of Southern blots. See Am. J. Hum.Genet. p32, 314-331 (1980) or Sci. Am. 258, 40-48 (1988). Since mutations often occur randomly they may affect the recognition sequence of the endonuclease and preclude the enzymatic cleavage at that site. Restriction fragment length polymorphism mappings (RFLPS) are based on changes at the restriction site. They are accurate but not very informative (PIC› 0.3). The major problem with RFLPs is the inability of a test to detect changes that do not affect cleavage with a restriction endonuclease. In addition, the methods used to detect RFLPs are very labor intensive and expensive, especially the techniques which includes Southern blot analysis.

[0008] Another technique for detecting specific mutations in particular DNA segment involves hybridizing DNA segments which are being analyzed with a complementary, labeled oligonucleotide probe. See Nucl. Acids Res. 9, 879-894 (1981). Since DNA duplexes containing even a single base pair mismatch exhibit high thermal instability, the differential melting temperature can be used to

distinguish target DNAs that are perfectly complimentary to the probe from target DNAs that only differ by a single nucleotide. See, *e.g.,* U.S. Pat. No. 4,683,194. Further, subtle genetic differences among related individuals regarding nucleotides which are substituted in the DNA chains are difficult to detect. VNTR's or Jeffrey's probes are very informative but labor intensive, in distinction to microsatellites which are equally informative PCR based tests.

[0009] Short tandem repeat (STR) polymorphisms are commonly used in DNA identification, either as adjuncts to other genetic tests, or as stand-alone tests. Typically, when STRs are used for human identification, they are amplified in groups of three to four loci (multiplex amplification). Generally, the resulting amplified fragments are analyzed by polyacrylamide gel electrophoresis. Polymorphisms are thus typed according to size by comparing to similarly labeled known external standards or differently labeled internal standards. U.S. Pat. No. 5,364,759 describes the genus of simple tandem repeats as well as a DNA typing method employing the simple tandem repeats and PCR amplification of the loci. Fragments are analyzed by differential labeling of the products.

[0010] A critical parameter in DNA typing is the power of exclusion for the system. Power of exclusion is the ability of a test to exclude a falsely accused individual based on the individual's genetic characteristics. The commonly used STR multiplexes have exclusion probabilities in the range of 85% to 91%. This compares unfavorably with restriction fragment length polymorphic loci (RFLP loci), which often provide an equivalent power with just one locus. STR testing batteries which include greater numbers of lower power systems are more susceptible to this problem than are RFLP testing batteries which include a smaller number of higher power systems. The low exclusion probabilities of commonly used STR loci are the most negative aspect of their use, although the frequencies of both alleles of an individual can be included in calculating match. Although it is simpler and faster to perform DNA typing with STR loci than with RFLP loci and it can be performed with much smaller quantities of DNA, typing using STR loci sacrifice in exclusion power. Another disadvantage of current STR multiplex DNA typing systems is that the amplification is rarely, if ever, clean. In

other words there is considerable formation of spurious bands, which is thought to be due to DNA polymerase slippage and mis-priming events (see e.g., Tautz D., Hyper variability of Simple Sequences as a General Source for Polymorphic DNA Markers, Nuc. Acids Res., 17(16) 6463-70 (1989)).

[0011] Other polymorphisms take the form of single nucleotide variations between individuals of the same species. Such polymorphisms are far more frequent than RFLPS, STRs and VNTRs. Some single nucleotide polymorphisms occur in protein-coding sequences, in which case, one of the polymorphic forms may give rise to the expression of a defective or other variant protein and, potentially, a genetic disease. Other single nucleotide polymorphisms occur in noncoding regions. Some of these polymorphisms may also result in defective protein expression (e.g., as a result of defective splicing). Other single nucleotide polymorphisms have no phenotypic effects.

[0012] Single nucleotide polymorphisms (SNPs) can be used in the same manner as RFLPs, and VNTRs but offer several advantages. Single nucleotide polymorphisms occur with greater frequency and are spaced more uniformly throughout the genome than other forms of polymorphism. The greater frequency and uniformity of single nucleotide polymorphisms means that there is a greater probability that such a polymorphism will be found in close proximity to a genetic locus of interest than would be the case for other polymorphisms. Also, the different forms of characterized single nucleotide polymorphisms are often easier to distinguish than other types of polymorphism, *e.g.*, by use of assays employing allele-specific hybridization probes or primers).

[0013] There is a need in the art for a very accurate genetic relationship test procedure which uses very small amounts of an original DNA sample, yet produces very accurate results. This is particularly true in the forensic medicine area and criminology because often only very small samples of DNA available.

SUMMARY OF THE INVENTION

[0014] The present invention provides novel polymorphisms on the Y chromosome and methods of using Y chromosome polymorphisms as indicators of evolutionary heritage. The polymorphisms of particular interest in the present invention are clustered to specific regions of the Y chromosome, with polymorphisms of particular use found mostly in the Non-recombining Region of the human Y chromosome (NRY). These polymorphisms, including but not limited to SNPs, insertions, and deletions, may be useful for numerous applications, including forensics, paternity testing, diagnosis and the like.

[0015] In one embodiment, the present invention provides nucleic acid segments of between 10 and 100 bases containing at least 10, 15 or 20 contiguous nucleotides from any of the polymorphic regions of the Y chromosome shown in TABLE 1, and may include a polymorphic site. Complements of these segments are also included. The segments can be DNA or RNA, and can be double or single-stranded. Some segments are 10-20 or 10-50 bases long and may be less than 20 or 50 bases long. Preferred nucleic acid segments allow for the identification and analysis of nucleic acid sequences on the Y chromosome which include at least one polymorphic site that is at least diallelic.

[0016] The invention further provides allele-specific oligonucleotides that hybridize to a polymorphic region marker (M1 to M319 (excluding unassigned markers) of the Y chromosome as shown in TABLE 1, or its complement. These oligonucleotides can be probes or primers. In a particular embodiment, the nucleic acid segments include the forward and/or reverse primer sequences (e.g. primer pairs) as in Table 1. Primer pairs allow for the amplification and identification of specific polymorphic regions of the Y chromosome. Polymorphic regions of interest for amplification and/or identification include but are not limited to the NRY regions of the Y chromosome. The polymorphic regions (polymorphic markers) shown in TABLE 1 are nucleic acids of about between 100 and 700 bases, about 200 to about 600 bases and, in some embodiments, about 250 to about 500 bases in length. Many of the polymorphic nucleic acids (polymorphic

regions (markers) shown in TABLE 1 may include more than one polypmorphic site.

[0017] The invention further provides a method of analyzing a nucleic acid from an individual. The method determines which base is present at any one of the polymorphic sites of the Y chromosome as shown in TABLE 1 in bold type. Optionally, a set of bases occupying a set of the polymorphic sites shown in TABLE 1 is determined. This type of analysis can be performed on a plurality of individuals who are tested for the presence of a particular polymorphism by identifying specific polymorphic markers. The polymorphism can be correlated with a base or set of bases present at the polymorphic sites in the individuals tested, and the evolutionary heritage of the individual can be indicated by the presence or absence of a particular polymorphism.

[0018] In one embodiment, the invention provides a method for determining the ethnic origin of a male, comprising obtaining a nucleic acid sample from the male and identifying at least two polymorphic markers in the nucleic acid sample indicative of the ethnic origin of the male, using at least one primer pair from TABLE 1. The identifying of the polymorphic markers may indicate the ethnic origin of the male as being at least one of the haplotype groups selected from the group consisting of haplotype Group I, Group II, Group III, Group IV, Group V, Group VI, Group VII, Group VIII, Group IX or Group X. In some embodiments, at least one polymorphic marker identified is a polymorphic marker from TABLE 1. The polymorphic markers may identify a haplotype associated with a haplotype group selected from the group consisting of haplotype Group I, Group II, Group III, Group IV, Group V, Group VI, Group VII, Group VIII, Group IX or Group X, or a sub-haplotype group for the ethnic origin of the male.

[0019] In another embodiment, the invention provides a method for identifying a plurality of polymorphic sites in a nucleic acid, comprising obtaining a sample of the nucleic acid from at least one individual, and identifying, in the nucleic acid, at least one of the polymorphic sites in at least two polymorphic markers of

TABLE 1. The sample of nucleic acids may be obtained from a plurality of individuals, with the presence of the polymorphic markers in each sample of the nucleic acid determined for each of the individuals. The method may further comprise testing each individual for presence of a group of polymorphic markers which identify the haplotype of each individual, wherein the haplotype is indicative of a geographic distribution of a population or an ancestral population.

[0020] In still other embodiments, the invention provides a method for determining the ethnic origin of a human male individual, comprising obtaining a nucleic acid sample from the male, testing the nucleic acid sample for presence of a plurality of polymorphic markers selected from TABLE 1, identifying which polymorphic markers are present in the nucleic acid sample, and assigning a haplotype group to the male based on the identified markers, wherein the haplotype group is indicative of the ethnic origin of the male.

[0021] In certain embodiments, the invention provides a method for determining the paternity of a human male individual, comprising obtaining a nucleic acid sample from the male, testing the nucleic acid sample for the presence of a plurality of polymorphic markers from TABLE 1, identifying which polymorphic markers are present in the nucleic acid sample, and comparing the identified polymorphic markers to a set of polymorphic markers identified in nucleic acid samples from potential fathers.

[0022] The invention additionally provides a kit for determining ethnic origin of an individual, comprising at least two primer pairs capable of identifying at least two polymorphic markers from TABLE 1. The kit may further comprise a control nucleic acid for detecting the presence or absence of the polymorphic markers from TABLE 1.

[0023] The invention further comprises a set of primers and enzymes useful in performing an assay to identify particular polymorphisms in human male DNA.

A method of identifying polymorphisms is disclosed whereby a sample is provided and subjected to amplification using primers of the invention and thereafter determining sequences (polymorphic regions) which were amplified.

[0024] A feature of the invention is that polymorphisms not previously identified are described herein, and are associated with a particular haplotype, indicative of a specific evolutionary heritage.

[0025] An advantage of the invention is that the sequences disclosed herein can be used in a range of different assay systems to determine the presence of a polymorphism in a sample.

[0026] A feature of the invention is a method for analyzing a set of unique polymorphisms on the Y chromosome to determine and identify an individual's evolutionary heritage and/or ethnicity.

[0027] A feature of the invention is to provide a kit for determining an individual's geographical or ethnic origins.

[0028] These and other objects, advantages, and features of the invention will become apparent to those persons skilled in the art upon reading the details of the invention as fully described below.

BRIEF DESCRIPTION OF THE DRAWINGS

[0029] Fig. 1. Contemporary worldwide distribution of Y chromosome groups in 22 regions determined by the methods and compositions of the invention.

[0030] **Fig. 2**. A phylogenetic tree deduced from 167 NRY polymorphisms on the principle of maximum parsimony.

[0031] **Fig. 3.** Maximum likelihood network inferred from the haplotype frequencies.

[0032] **Fig. 4.** Maximum parsimony phylogeny of human NRY chromosome biallelic variation.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

[0033] Before the present polymorphisms and detection methods are described, it is to be understood that this invention is not limited to particular methods or polymorphisms described, as such may, of course, vary. It is also to be understood that the terminology used herein is for the purpose of describing particular embodiments only, and is not intended to be limiting, since the scope of the present invention will be limited only by the appended claims.

[0034] Where a range of values is provided, it is understood that each intervening value, to the tenth of the unit of the lower limit unless the context clearly dictates otherwise, between the upper and lower limits of that range is also specifically disclosed. Each smaller range between any stated value or intervening value in a stated range and any other stated or intervening value in that stated range is encompassed within the invention. The upper and lower limits of these smaller ranges may independently be included or excluded in the range, and each range where either, neither or both limits are included in the smaller ranges is also encompassed within the invention, subject to any specifically excluded limit in the stated range. Where the stated range includes one or both of the limits, ranges excluding either or both of those included limits are also included in the invention.

[0035] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Although any methods and materials similar or equivalent to those described herein can be used in the practice or testing of the present invention, the preferred methods and materials are now described. All publications mentioned herein are incorporated herein by reference to disclose and describe the methods and/or materials in connection with which the publications are cited.

[0036] It must be noted that as used herein and in the appended claims, the singular forms "a", "and", and "the" include plural referents unless the context

clearly dictates otherwise. Thus, for example, reference to "a nucleic acid" includes a plurality of such nucleic acids and reference to "the primer" includes reference to one or more primers and equivalents thereof known to those skilled in the art, and so forth.

[0037] The publications discussed herein are provided solely for their disclosure prior to the filing date of the present application. Nothing herein is to be construed as an admission that the present invention is not entitled to antedate such publication by virtue of prior invention. Further, the dates of publication provided may be different from the actual publication dates which may need to be independently confirmed.

THE INVENTION IN GENERAL

[0038] The use of certain nucleotide repeat polymorphisms for identifying or comparing DNA segments have been described. (See *e.g.*, Weber & May *Am Hum Genet* 44:388 (1989), Litt & Luthy *Am Hum Genet* 44:397(1989)). The present invention is based on the finding that particular polymorphisms on the Y chromosome, including the novel polymorphisms included herein, are indicative of the evolutionary heritage and/or a paternal lineage in an individual having a Y chromosome (e.g., a male or XXY individual). These particular polymorphic genetic segments, and primers used to identify the polymorphisms for identification and comparison purposes, correspond to regions of the Y chromosome having clustered polymorphisms that are homopolymeric in regions which exhibit a very low mutation rate. An advantage of the polymorphisms of the invention is that no recombination occurs in the regions containing these markers, and thus the accumulation of mutations is preserved as an intact haplotype. This creates a genetic profile that remains intact across the generations. If men share the same derived allele, then they are identical by descent, not just by state. While a very small amount of recurrent or revertant back mutation has been observed at some markers, these anomalies are easily recognized as such because of the high resolution of the Y tree. The recognition

of new Y-chromosome markers represents a major leap in the investigation of human genetic diversity (in male lineages, complementing the information from female lineages derived from mitochondrial DNA).

[0039] The polymorphisms and methods of the present invention provide a simple way of identifying male siblingship as well as a genetic route to identify male children by so called "genebanking" using DNA or blood, or saliva from a child. Also the Y chromosome polymorphisms can reveal patterns (estimates) of recent gene flow from one gene pool to another, i.e. admixture. The methods of the present invention make the large amount of information contained in the phylogeny of haplotypes accessible for analysis.

DEFINITIONS

[0040] The term "oligonucleotide" as used herein can be DNA, RNA, or a substituted variation of these nucleic acids. The oligonucleotide may be single- or double-stranded. Oligonucleotides can be naturally occurring or synthetic, but are typically prepared by synthetic means. Preferred oligonucleotides of the invention include segments of DNA, or their complements including any one of the polymorphic sites shown in TABLE 1. The segments are usually between 5 and 100 bases (nucleotides), and often between 5-10, 5-20, 10-20, 10-50, 20-50 or 20-100 bases. The polymorphic site can occur within any position of the segment. The segments can be from any of the allelic forms of DNA shown in TABLE 1.

[0041] The term "hybridization probes" as used herein refers to oligonucleotides capable of binding in a base-specific manner to a complementary strand of nucleic acid. Such probes include peptide nucleic acids, as described in Nielsen et al., Science 254, 1497-1500 (1991).

[0042] The term "primer" as used herein refers to an oligonucleotide having at least a single-stranded portion that is adapted to act as a point of initiation of template-directed DNA synthesis under appropriate conditions (i.e., in the presence of four different nucleoside triphosphates and an agent for polymerization, such as, DNA or RNA polymerase or reverse transcriptase) in an

appropriate buffer and at a suitable temperature. The appropriate length of a primer depends on the intended use of the primer but typically ranges from 15 to 30 nucleotides. A primer need not reflect the exact sequence of the template but must be sufficiently complementary to hybridize with a template.

[0043] The term "primer site" as used herein refers to the area of the target DNA to which a primer hybridizes. The term "primer pair" as used herein refers to a set of primers including at least one 5' upstream primer that hybridizes with the 5' end of the DNA sequence to be amplified (a forward or "for" primer) and at least one 3' downstream primer that hybridizes with the complement of the 3' end of the sequence to be amplified (a reverse or "rev" primer). Primer pairs allow for the amplification and identification of corresponding polymorphic regions.

[0044] The term "polymorphic site" is used herein to describe mutations within a nucleic acid sequence which include but are not limited to site specific mutations, insertions and deletions, these mutations being found in the nucleic acid of some individuals and not in others, e.g. the polymorphic site identifies a specific polymorphism of an individual. The present invention provides segments of nucleic acid which contain at least one polymorphic site (i.e. polymorphic region). These "polymorphic regions" of the Y chromosome can be analyzed to identify a specific polymorphic site which in turn identifies a specific polymorphism associated with certain individuals.

[0045] The polymorphic regions of the present invention are also defined as "polymorphic markers" due to their usefulness in marking (identifying specific polymorphic sites). The polymorphic markers of the present invention identify specific haplotypes in the male population, these haplotypes being indicative of a specific geographical or ethnic origin. Certain polymorphic markers which identify a polymorphism shared by a large group of individuals allow for the grouping of those haplotypes which share that marker. These more commonly found markers are found at the branch points of a phylogenetic tree and are crucial in separating individuals into unique haplotype groups. The haplotype groups have this ancestral marker which branches off from a point earlier in the

phylogenetic tree. The polymorphic markers of the present invention have identified over 171 haplotypes which can be divided into ten haplotype groups.

[0046] The term "polymorphism" as used herein refers to the occurrence of two or more genetically determined alternative sequences or alleles in a population. A polymorphic marker or site is the locus at which divergence occurs. Preferred markers have at least two alleles, each occurring at a frequency of greater than 1%, and more preferably greater than 10% or 20% of a selected population, and can be present at a frequency greater than 30% to 50% or more in selected portions of the population. A polymorphic locus may be as small as one base pair. Polymorphic markers include restriction fragment length polymorphisms, VNTR's, hypervariable regions, minisatellites, dinucleotide repeats, trinucleotide repeats, tetranucleotide repeats, simple sequence repeats, and insertion elements such as Alu. The first identified allelic form is arbitrarily designated as the reference form and other allelic forms are designated as alternative or variant alleles. Polymorphisms refer to sequence differences between a reference form and a selected allele, and encompasses single or multiple nucleotide differences which can result from nucleotide insertion(s), deletion(s), substitution(s) and/ or a combination thereof. The allelic form occurring most frequently in a selected population is sometimes referred to as the wildtype form. Diploid organisms may be homozygous or heterozygous for allelic forms. A diallelic polymorphism has two forms. A triallelic polymorphism has three forms. The term "polymorphism" as used herein refers to any detectable polymorphic site in DNA or RNA that is detectable using the present methods. The term as used herein encompasses, for example, polymorphisms associated with a disease state (i.e. mutations), "silent" polymorphisms (i.e. associated with a wild-type phenotype or in a non-coding region), and polymorphisms associated with a predisposition and/or response to treatment (i.e. a polymorphism in an allele of a gene).

[0047] The term "single nucleotide polymorphism" and "SNP" as used interchangeably herein refers to a polymorphic site occupied by a single nucleotide (i.e. single base), which is the site of variation between allelic

sequences. In general, SNPs are DNA sequence variations that occur when a single nucleotide (A, T, C or G) in the genomic sequence is altered. For example a SNP might change the DNA sequence AAGGCTAA to ATGGCTAA. SNPs can occur in both coding (gene) and noncoding regions of the genome. The site is usually preceded by and followed by highly conserved sequences of the allele (*e.g.,* sequences that vary in less than 1/100 or 1/1000 members of the population).

[0048] A single nucleotide polymorphism usually arises due to substitution of one nucleotide for another at the polymorphic site. A transition is the replacement of one purine by another purine or one pyrimidine by another pyrimidine. A transversion is the replacement of a purine by a pyrimidine or vice versa. Single nucleotide polymorphisms can also arise from a deletion of a nucleotide or an insertion of a nucleotide relative to a reference allele. Hybridizations are usually performed under stringent conditions, for example, at a salt concentration of no more than 1M and a temperature of at least 25°C. For example, conditions of 5X SSPE (750 mM NaCl, 50 mM NaPhosphate, 5 mM EDTA, pH 7.4) and a temperature of 25°-30°C are suitable for allele-specific probe hybridizations.

[0049] The term "isolated nucleic acid" as used herein refers to a nucleic acid isolated from an individual that is the predominant species present (i.e., on a molar basis it is more abundant than any other individual species in the composition). Preferably, an isolated nucleic acid comprises at least about 50, 80 or 90 percent (on a molar basis) of all macromolecular species present. Most preferably, the object species is purified to essential homogeneity, i.e. contaminant species cannot be detected in the composition by conventional detection methods. The isolated nucleic acid includes a selected DNA fragment (*e.g.,* isolated by an amplification reaction), and an isolated mRNA.

[0050] The term "evolutionary heritage" as used herein refers to the association of a particular polymorphism with a population having a particular geographic distribution. This includes polymorphisms that are indicative of an ancestral population, i.e. a population from which an individual is a descendant.

GENERAL ASPECTS OF THE INVENTION

[0051] The present application provides novel polymorphisms, including polymorphisms clustered in and around a non-recombining portion of the human Y chromosome (NRY) The polymorphic sites and the regions flanking these polymorphic sites are shown in TABLE 1.

[0052] By knowing sequences which include particular polymorphisms on the Y chromosome, primers based on these sequences can be used in detection assays. The primers can be provided in assay kits which cover from one to any and all of the polymorphisms developed here and the kits may further comprise appropriate enzymes for use with the primers and/or reagents for the isolation and processing of nucleic acids from an individual.

[0053] The methods and compositions of the present invention allow for the genetic typing of male individuals into ten major haplotype groups. The markers and primer sets shown in TABLE 1 allow not only for typing males into one of the haplotype groups or a combination of haplotype groups, but also enables an individual to be identified to a specific geographical area associated with haplotype group. Figure 1 shows a contemporary worldwide frequency distribution of the 10 Y chromosome groups in 22 regions. Each group is represented by a distinguishing color. Colored sectors reflect representative group frequencies. The frequency distribution of the ten groups is based on > 1000 globally diverse samples genotyped using a hierarchical top down approach as illustrated in FIG.1 above the global map. The representative branching and frequency of polymorphic markers in TABLE 1 are also shown in FIG. 1 (individual marker numbers are not shown).

[0054] The identification of an individuals haplotype is base on identifying the presence of at least two distinct polymorphic markers (i.e. at least two distinct polymorphic sites must be identified), for example, polymorphic markers M91 and M278 identify haplotype 9(shown in FIG. 2 and FIG. 4). More likely, determining the haplotype of an individual involves the identification of 3 or more

markers, usually at least about 3 to 7 markers, or 7 to 9 markers or even 9 or more markers.

[0055] Haplotype groups comprise haplotypes which have at least one ancestral marker which branches off from a point earlier in the phylogenetic tree. For example, marker 91 (M91) identifies haplotypes in Group I while haplotypes in group V are identified by one marker from each of the following sets of markers; one marker from {M42, M94, M139, M251, M299} plus one from {M168, M294} and one marker from {RPS4Y, M216, M316}. To determine which haplotype group and individual is associated with, the individuals nucleic acid would need to be analyzed with at least eleven polymorphic markers. For exemplary purposes, an individuals nucleic acid could be assayed for the presence and absence of the following markers; M91, M299, M249, M294, M203, M96, M316, M9, M74, M207, M214 to determine which haplotype group they are associated with which is indicative of a certain geographical or ethnic origin.

[0056] Fig. 1 illustrates that haplotype Group I is mainly associated with Africa and in particular, southern and eastern Africa (approximately about 90% of males of haplotype Group I are of African origin). Haplotype Groups II (about 80% to about 99% frequency distribution (f.d.)) and III (about 75% to about 95% f.d.) are also strongly related to Africa compared to Groups IV through X. Populations represented in Groups I and II include some Khoisan and Bantu speakers from South Africa, Pygmies from central Africa, and lineages in Sudan, Ethiopia and Mali. Virtually all men with Group I and II haplotypes are of African affiliation from a paternal perspective. Group III lineages are predominantly African, although a sub-set of Group III lineages occur in populations bordering the Mediterranean (Middle East, Turkey, North Africa, Southern Europe).

[0057] Approximately about 70% to about 99% of the males in Group IV are of Japanese origin. Group V is slightly associated with Japan (about 10% to about 25% f.d.) and Indonesia (about 10% to about 35% frequency) with the largest frequency being associated with Australia and central Asians (about 45% to about 75% f.d.).

[0058] Group VI is more widely distributed than other haplotypes, covering the geographical area of Europe, Eastern Europe, Asia, and India. The presence of haplotype group VI in North America, Australia and Polynesia is a consequence of recent human movements since C. Columbus catalyzed the age of exploration. The largest Group VI frequency is associated with southern Europe and the middle east, with a distribution frequency of about 60% to about 85%.

[0059] Group VII is more widely associated with eastern Asia and Indonesia with distribution frequencies ranging from about 75% to about 99%. Group VIII is almost exclusively found in Papua-New Guniea (distribution frequencies of about 70% to about 95%) with a slight distribution in central Asia (distribution frequency of about 1% to about 30%). Recently, there is evidence which indicates the presence of Group VIII in Indonesia. Other specific Group VIII lineages occur in India and Europe. Individuals of haplotype Group IX are mostly associated Europe (about 75% to about 95% f.d.), India (about 25% to about 50% f.d.). Their occurrence in North America (about 35% to about 55%) Australia (35%), Polynesia is a consequence of European gene flow during the last 500 years.

[0060] Group X individuals are geographically associated with Central Asia and the Americas with a frequency distribution in North America of about 25% to about 50%, Central America of about 75% to about 95% and in South America of about 80% to about 99%. The above distribution frequencies of the various haplotypes in the geographic regions mentioned above are only representative ranges of the haplotype frequencies worldwide.

Analysis of Polymorphisms

[0061] Polymorphisms are detected in a target nucleic acid from an individual being analyzed. For assay of genomic DNA, virtually any biological sample (other than pure red blood cells) is suitable. For example, convenient tissue samples include whole blood, semen, saliva, tears, urine, fecal material, sweat, buccal, skin and hair. For assay of cDNA or mRNA, the tissue sample must be obtained from an organ in which the target nucleic acid is expressed. For

purposes of the present invention, the sample is obtained from a male, and preferably a human male.

[0062] Many of the methods described below require amplification of DNA from target samples. This can be accomplished by *e.g.,* PCR. See generally PCR Technology: Principles and Applications for DNA Amplification (ed. H. A. Erlich, Freeman Press, N.Y., N.Y., 1992); PCR Protocols: A Guide to Methods and Applications (eds. Innis, et al., Academic Press, San Diego, Calif., 1990); Mattila et al., Nucleic Acids Res. 19, 4967 (1991); Eckert et al., PCR Methods and Applications 1, 17 (1991); PCR (eds. McPherson et al., IRL Press, Oxford); and U.S. Pat. No. 4,683,202.

[0063] Other suitable amplification methods include the ligase chain reaction (LCR) (see Wu and Wallace, Genomics 4, 560 (1989), Landegren et al., Science 241, 1077 (1988), transcription amplification (Kwoh et al., Proc. Natl. Acad. Sci. USA 86, 1173 (1989)), and self-sustained sequence replication (Guatelli et al., Proc. Nat. Acad. Sci. USA, 87, 1874 (1990)) and nucleic acid based sequence amplification (NASBA). The latter two amplification methods involve isothermal reactions based on isothermal transcription, which produce both single stranded RNA (ssRNA) and double stranded DNA (dsDNA) as the amplification products in a ratio of about 30 or 100 to 1, respectively.

Detection of Polymorphisms in Target DNA

[0064] There are two distinct types of analysis depending whether a polymorphism in question has already been characterized. The first type of analysis is sometimes referred to as *de novo* characterization. This analysis compares target sequences in different individuals to identify points of variation, e.g., polymorphic sites, SNPs. By analyzing groups of individuals representing the greatest ethnic diversity among humans and greatest breed and species variety in plants and animals, patterns characteristic of the most common alleles/haplotypes of the locus can be identified, and the frequencies of such populations in the population determined. Additional allelic frequencies can be determined for subpopulations characterized by criteria such as geographical

distribution and ancestral ethnicity. The *de novo* identification of the polymorphisms of the invention is described in the Examples section. The second type of analysis is determining which form(s) of a characterized polymorphism are present in individuals under test. There are a variety of suitable procedures, which are discussed in turn.

Allele-Specific Probes

[0065] The design and use of allele-specific probes for analyzing polymorphisms is described by e.g., Saiki et al., Nature 324, 163-166 (1986); Dattagupta, EP 235,726, Saiki, WO 89/11548. Allele-specific probes can be designed that hybridize to a segment of target DNA from one individual but do not hybridize to the corresponding segment from another individual due to the presence of different polymorphic forms in the respective segments from the two individuals. Hybridization conditions should be sufficiently stringent that there is a significant difference in hybridization intensity between alleles, and preferably an essentially binary response, whereby a probe hybridizes to only one of the alleles. Probes with such specificity allow for the determination of a specific base occupying a polymorphic site in a sequence of a polymorphic region. Some probes are designed to hybridize to a segment of target DNA such that the polymorphic site aligns with a central position (e.g., in a 15 mer at the 7 position; in a 16 mer, at either the 8 or 9 position) of the probe. This design of probe achieves good discrimination in hybridization between different allelic forms.

[0066] Allele-specific probes are often used in pairs, one member of a pair showing a perfect match to a reference form of a target sequence and the other member showing a perfect match to a variant form. Several pairs of probes can then be immobilized on the same support for simultaneous analysis of multiple polymorphisms within the same target sequence.

Tiling Arrays

[0067] The polymorphisms can also be identified by hybridization to nucleic acid arrays, some example of which are described by WO 95/11995. The same array

or a different array can be used for analysis of characterized polymorphisms. WO 95/11995 also describes subarrays that are optimized for detection of a variant form of a precharacterized polymorphism. Such a subarray contains probes designed to be complementary to a second reference sequence, which is an allelic variant of the first reference sequence. The second group of probes is designed by the same principles as described in the Examples except that the probes exhibit complementarily to the second reference sequence. The inclusion of a second group (or further groups) can be particular useful for analyzing short subsequences of the primary reference sequence in which multiple mutations are expected to occur within a short distance commensurate with the length of the probes (i.e., two or more mutations within 9 to 21 bases).

Allele-Specific Primers

[0068] An allele-specific primer hybridizes to a site on target DNA overlapping a polymorphism and only primes amplification of an allelic form to which the primer exhibits perfect complementarily. See Gibbs, Nucleic Acid Res. 17, 2427-2448 (1989). This primer is used in conjunction with a second primer which hybridizes at a distal site. Amplification proceeds from the two primers leading to a detectable product signifying the particular allelic form is present. A control is usually performed with a second pair of primers, one of which shows a single base mismatch at the polymorphic site and the other of which exhibits perfect complementarily to a distal site. The single-base mismatch prevents amplification and no detectable product is formed. The method works best when the mismatch is included in the 3'-most position of the oligonucleotide aligned with the polymorphism because this position is most destabilizing to elongation from the primer. See, e.g., WO 93/22456.

Direct-Sequencing

[0069] The direct analysis of the sequence of polymorphisms of the present invention can be accomplished using either the dideoxy chain termination method or the Maxam Gilbert method (see Sambrook et al., Molecular Cloning, A

Laboratory Manual (2nd Ed., CSHP, New York 1989); Zyskind et al., Recombinant DNA Laboratory Manual, (Acad. Press, 1988)). In a preferred embodiment, the direct sequencing would be carried using fluorescent sequencing, *e.g.,* using a PE Biosystems 373A sequencer.

Denaturing Gradient Gel Electrophoresis

[0070] Amplification products generated using the polymerase chain reaction can be analyzed by the use of denaturing gradient gel electrophoresis. Different alleles can be identified based on the different sequence-dependent melting properties and electrophoretic migration of DNA in solution. Erlich, ed., PCR Technology, Principles and Applications for DNA Amplification, (W.H. Freeman and Co, New York, 1992), Chapter 7.

Single-Strand Conformation Polymorphism Analysis

[0071] Alleles of target sequences can be differentiated using single-strand conformation polymorphism analysis, which identifies base differences by alteration in electrophoretic migration of single stranded PCR products, as described in Orita et al., Proc. Nat. Acad. Sci. 86, 2766-2770 (1989). Amplified PCR products can be generated as described above, and heated or otherwise denatured, to form single stranded amplification products. Single-stranded nucleic acids may refold or form secondary structures which are partially dependent on the base sequence. The different electrophoretic mobilities of single-stranded amplification products can be related to base-sequence difference between alleles of target sequences.

Detection of SNP Polymorphisms

[0072] Where the polymorphism is a SNP, any suitable method known in the art can be used in their detection. For example, the present methods can utilize the detection of SNPs by DHPLC (see U.S. Pat. No. 5,795,976) to isolate and analyze specific SNPs on the Y chromosome of a large number of individuals in a fast, efficient and inexpensive manner. This method involves separating heteroduplex

and homoduplex nucleic acid molecules (e.g., DNA or RNA) in a mixture using high performance liquid chromatography under partially denaturing conditions. In a preferred embodiment, the SNPs are identified on the Y chromosome using techniques such as those disclosed in co-pending application US Application Serial No. 09/502,558, February 10, 2000.

Mass Spectrometry

[0073] Mass spectrometry can also be used in the methods of the present invention to verify a polymorphism and/or to identify additional polymorphisms. The mass spectrum of a nucleic acid containing the polymorphic site can be compared to the mass spectrum of nucleic acids obtained from samples of known residues at the polymorphic site. These known spectra are referred to as "signature" spectra. A simple comparison of the sample spectrum vs. signature spectra will reveal whether an individual's DNA has a specific base occupying the polymorphic site. Although sequencing of fragments of nucleic acids is possible using mass spectrometry, actual sequencing of the nucleic acid is not required for this mutational analysis. Less preparation and analysis is needed to prepare and analyze a complete, intact fragment as compared to treating a sample for actual sequencing.

[0074] Certain mass spectrometry techniques can be used to analyze for polymorphisms. Short oligomers, *e.g.,* from one nucleotide up to approximately 50 nucleotides, can be analyzed and the resulting spectra compared with signature spectra of samples known to be wild-type or to contain a known polymorphism. A comparison of the locations (mass) and heights (relative amounts) of peaks in the sample with the known signature spectra indicate what type of polymorphism, if any, is present. Exemplary protocols are described in U.S. Pat Nos. 5,872,003, 5,869,242, 5,851,765, 5,622,824, and 5,605, 798, which are incorporated herein by reference for teaching such techniques.

[0075] After determining polymorphic form(s) present in an individual at one or more polymorphic site on the Y chromosome, this information can be used in a number of methods.

Methods of Use of the Polymorphisms of the Invention

[0076] The methods of the invention have utility in a wide variety of fields where it is desirable to identify known polymorphisms of a particular individual and/or to determine allelic distribution in a group or population. Such methods include, but are not limited to, linkage analysis for the identification of disease loci, evolutionary studies to determine rates of evolution in a population, identification of polymorphisms useful in forensic identification, identification of mutations associated with a disease or predisposition, genetic marker development, and the like.

Forensics

[0077] Determination of which polymorphic sites an individual possesses, identifies a haplotype, which refers to a set of polymorphic markers that distinguishes the individual. See generally National Research Council, *The Evaluation of Forensic DNA Evidence* (Eds. Pollard et al., National Academy Press, DC, 1996). Since the polymorphic sites of the invention are generally within a region of about 50,000 bp in the human genome, the probability of recombination between these polymorphic sites is low. The more sites that are analyzed the lower the probability that the set of polymorphic markers for one individual is the same as that in an unrelated individual. If multiple polymorphic sites are analyzed, the sites are usually in different polymorphic regions (on different polymorphic markers). Thus, polymorphisms of the invention may be used in conjunction with polymorphisms in distal genes. Preferred polymorphisms for use in forensics are diallelic because the population frequencies of two polymorphic forms can usually be determined with greater accuracy than those of multiple polymorphic forms at multi-allelic loci.

[0078] An exemplary set of polymorphic markers useful for identifying the haplotype group of an individual are the following; Markers 304(Group VI, Mediterranean), 242 (Group X, C. Asia, India, Americas), 269 (Group IX, W. Europe), 207 (Group IX, Europe, W. Asia), 74 (Groups IX-X, global), 214 (Group VII, E. Asia), 9 (Groups VII-X, global), 235 (Groups VI-X, global), 316 (Group V, Asia, America, Polynesia, Melanesia), 174 (Group IV, Asia, Japan), 299 (Groups II-X, global), 246 (Group I, Africa), 249 (Group II, Africa) 294 (Groups III-X, global), 96 (Group III, Africa, Mediterranean).

[0079] The capacity to identify a distinguishing or unique set of forensic markers in an individual is useful for forensic analysis. For example, one can determine whether a blood sample from a suspect matches a blood or other tissue sample from a crime scene by determining whether the set of polymorphic forms occupying selected polymorphic sites is the same in the suspect and the sample. If the set of polymorphic markers does not match between a suspect and a sample, it can be concluded (barring experimental error) that the suspect was not the source of the sample. If the set of markers does match, one can conclude that the DNA from the suspect is consistent with that found at the crime scene. If frequencies of the polymorphic forms at the loci tested have been determined (e.g., by analysis of a suitable population of individuals), one can perform a statistical analysis to determine the probability that a match of suspect and crime scene sample would occur by chance. If several polymorphic loci are tested, the cumulative probability of non-identity for random individuals becomes very high (*e.g.*, one billion to one). Such probabilities can be taken into account together with other evidence in determining the innocence or guilt of an individual suspected of a criminal act.

[0080] The polymorphisms of the present invention are especially useful in identifying samples having genetic material from multiple individuals, since the polymorphisms are single copy. Thus, the detection of more than one polymorphic Y chromosome allele in a single sample is indicative of the presence of nucleic acids from multiple individuals within the sample. Such information can be useful, for example, when multiple perpetrators are suspected of

participating in a crime, or in the case of mixed unidentified remains at a grave site or accident scene.

[0081] The polymorphic sites and methods of the present invention are also useful in categorizing victims of violent crimes into ethnic and geographical groups. When a large number of victims need to be identified at a crime site, categorizing recovered victims by ethnicity can decrease the overall time for victim identification by reducing the number of comparison samples (samples from members of the victims family) to those of similar geographical origin.

Paternity Testing

[0082] The object of paternity testing is usually to determine whether a male is the father of a child. In most cases, the mother of the child is known and thus, the mother's contribution to the child's genotype can be traced. Paternity testing investigates whether the part of the child's genotype not attributable to the mother is consistent with that of the putative father. Paternity testing can be performed by analyzing sets of polymorphisms (polymorphic markers) in the putative father and the child. The polymorphic markers of the present invention can be useful in determining paternity of a male child, as they are specific to the Y chromosome. The mother need not be tested in such a case, as the mother has no contribution to the child's genotype as it pertains to the Y chromosome.

[0083] If the set of polymorphisms in the child attributable to the father does not match the putative father, it can be concluded, barring experimental error, that the putative father is not the real father. If the set of polymorphisms in the child attributable to the father does match the set of polymorphisms of the putative father, a statistical calculation can be performed to determine the probability of coincidental match. An exemplary method of determining the probability of parentage exclusion, i.e. the probability that a random male will have a polymorphic form at a given polymorphic site that makes him incompatible as the father) is described in WO 95/12607.

[0084] If several polymorphic loci are included in the analysis, the cumulative probability of exclusion of a random male is very high. This probability can be

taken into account in assessing the liability of a putative father whose polymorphic marker set matches the child's polymorphic marker set attributable to his father. This analysis can be further expanded to identify ancestral males (e.g., grandfather, great grandfather and so on). Such analysis can be useful in genealogical analysis, or in tracing the origin of ancestral man (e.g.) using samples obtained from an archeological site).

Longer-term Family Heritage

[0085] In addition to the use in paternity testing, the polymorphisms and methods of the present invention can be used to determine relationships through a paternal lineage for multiple generations. The constancy and low mutational rate of these regions of the Y chromosome allow an individual to trace his specific ancestral lineage using the Y chromosome polymorphisms. For example, a specific residue (base) in a polymorphic site may be indicative of a population that is in or from a certain region in Europe. Assaying an individual for this polymorphism can indicate that the individual's paternal ancestors were in or descended from this particular region.

Correlation of Polymorphisms with Phenotypic Traits

[0086] The polymorphisms of the invention may contribute to the phenotype of an organism in different ways. Some polymorphisms occur within a protein coding sequence and contribute to phenotype by affecting protein structure. The effect may be neutral, beneficial or detrimental, or both beneficial and detrimental, depending on the circumstances. Other polymorphisms occur in noncoding regions but may exert phenotypic effects indirectly via influence on replication, transcription, and translation.

[0087] A single polymorphism may affect more than one phenotypic trait. Likewise, a single phenotypic trait may be affected by polymorphisms in different genes. Further, some polymorphisms predispose an individual to a distinct mutation that is causally related to a certain phenotype.

[0088] Phenotypic traits include diseases that have known but hitherto unmapped genetic components. Phenotypic traits also include symptoms of, or susceptibility to, multifactorial diseases of which a component is or may be genetic, such as autoimmune diseases, inflammation, cancer, diseases of the nervous system, and infection by pathogenic microorganisms. Phenotypic traits also include characteristics such as longevity, appearance (e.g., baldness, obesity), strength, speed, endurance, fertility, and susceptibility or receptivity to particular drugs or therapeutic treatments.

[0089] Correlation is performed for a population of individuals who have been tested for the presence or absence of a phenotypic trait of interest and for polymorphic markers sets. To perform such analysis, the presence or absence of a set of polymorphisms (i.e. a polymorphic set) is determined for a set of the individuals, some of whom exhibit a particular trait, and some of which exhibit lack of the trait. The alleles of each polymorphism of the set are then reviewed to determine whether the presence or absence of a particular allele is associated with the trait of interest. Correlation can be performed by standard statistical methods such as a κ-squared test and statistically significant correlations between polymorphic form(s) and phenotypic characteristics are noted.

[0090] The polymorphisms and assays of the present invention are of particular use in determining the appropriate populations for mapping complex genetic traits and/or disorders. Population choice can be crucial for the success of gene mapping for particular traits and/or disorders. Populations having a high degree of inbreeding are also useful for linkage analysis (see, *e.g.,* Sheffield, VC et al., *Trends in Genetics* 4:391-6 (1998)), and the polymorphisms of the invention can be useful in determining the genetic heterogeneity of a population.

Antibodies to Specific Polymorphisms

[0091] Polyclonal and/or monoclonal antibodies that specifically bind to variant gene products but not to corresponding prototypical gene products are also provided. Antibodies can be made by injecting mice or other animals with the variant gene product or synthetic peptide fragments thereof. Monoclonal

antibodies are screened as are described, for example, in Harlow & Lane, Antibodies, A Laboratory Manual, Cold Spring Harbor Press, New York (1988); Goding, Monoclonal antibodies, Principles and Practice (2d ed.) Academic Press, New York (1986). Monoclonal antibodies are tested for specific immunoreactivity with a variant gene product and lack of immunoreactivity to the corresponding prototypical gene product. These antibodies are useful in diagnostic assays for detection of the variant form, or as an active ingredient in a pharmaceutical composition.

Use of the Present Method to Produce a Database of Y Chromosome Polymorphisms

[0092] The polymorphisms of the invention can be used as the basis for, or combined with other such polymorphisms to provide, a general catalog of genome variation to address the large-scale sampling designs required by association studies, gene mapping, and evolutionary biology. There is widespread interest in documenting the amount and geographic distribution of genetic variation in the human species. This information is desired by the biomedical community, whose work would be greatly facilitated by a densely packed map of polymorphic markers, particularly SNPs in the NRY region, to be used to for example, identify genes associated with disease by linkage disequilibrium between sets of adjacent markers and the occurrence of disease in populations, and to characterize disease-related variation among populations.

[0093] Anthropologists and archeologists use genetic variation to reconstruct our species' history, and to understand the role of culture and geography in the global distribution of human variation. The requirements for these two perspectives seem to be converging on a need for an accessible, representative DNA bank and statistical database of human variation.

[0094] In addition, these systems have potential in both routine forensic and intelligence database applications, either in place of or in conjunction with more traditional "DNA fingerprinting" databases produced using methods such as restriction fragment length polymorphism mapping.

[0095] The invention may be embodied in computer-readable media containing an electronically, magnetically, or optically stored code representative of the markers for polymorphic regions of Table 1, and/or stored code configured to create the electronically stored representation of Table 1 and the corresponding geographic distributions for these polymorphic markers (see TABLE 3). Such databases may be produced using a variety of different data configurations and processing capabilities. Examples include, but are not limited to, logical databases, physical databases, relational databases, central configuration databases, and the like. Database structures for genomic information may be based on, for example, the database structures disclosed in U.S. Patent No. 6,229,911. In other examples, the data generated for use in the present invention may be used to create a general database such as that described in U.S. Pat. No. 4,970,672 or a relational database such as that described in U.S. Pat. No. 5,884,311. Databases containing data generated for use in the methods of the invention may also be a central configuration database for data that is shared among multiprocessor computer systems. See U.S. Pat. No. 6,014,669. Other database systems and design methodologies can be found in I. Fogg and M. Orlowska, *Computers Math. Applic.* (UK), (1993) 25:97-106; S. Ceri, et al., Proceedings of the IEEE (1987) 75:533-545.

EXAMPLES

[0096] The following examples are put forth so as to provide those of ordinary skill in the art with a complete disclosure and description of how to make and use the present invention, and are not intended to limit the scope of what the inventors regard as their invention nor are they intended to represent that the experiments below are all or the only experiments performed. Efforts have been made to ensure accuracy with respect to numbers used (e.g. amounts, temperature, etc.) but some experimental errors and deviations should be accounted for. Unless indicated otherwise, parts are parts by weight, molecular weight is weight average molecular weight, temperature is in degrees Centigrade, and pressure is at or near atmospheric.

EXAMPLE 1

[0097] A phylogenetic tree was deduced from 167 polymorphisms from a Non-recombining Region of the human Y chromosome (NRY)on the principle of maximum parsimony (Figure 2). Seven of the 167 polymorphisms had been detected by means other than DHPLC and were taken from the literature to demonstrate the applicability of the method of the invention to polymorphisms with less demographic specificity than those in TABLE 1. Seventy-three of the 160 polymorphisms detected by DHPLC had been reported previously. Underhill, P. A. *et al Genome Res.* 7:996-1005 (1997). Shen, P. *et al Proc. Natl. Acad. Sci. USA* 97:7354-7359 (2000). Of the remaining 87 unreported polymorphisms, 53 were discovered in a set of 53 individuals of diverse geographic origin during the screening of the unique sequences and repeat elements, other than long interspersed elements, contained in three overlapping cosmid sequences (GenBank accession nos. AC003032, AC003095, AC003097) and a few small fragments scattered throughout the NRY. Finally, 34 were detected during genotyping. In total, the marker panel comprises 91 transitions, 53 transversions, 22 small insertions or deletions, and an *Alu* insertion. All polymorphisms are biallelic, except a double transversion, M116, that has three alleles, A, C or T, defining quite different haplotypes. Two non-CpG associated transitions (M64 and M108) showed evidence of recurrence but generated no ambiguities when considered in the context of other markers. The primer sequences used to detect the 167 polymorphisms are given in Table 1).

METHODS

[0098] **DNA samples.** The ascertainment set consisted of the following 53 samples with their subsequently determined haplogroup designations: *Africa*: 3 Central African Republic Biaka II, III (1); 2 Zaire Mbuti II, III; 2 Lissongo II, III; 2 Khoisan I, III; 1 Berta VI; 1 Surma I; 1 Mali Tuareg III; 1 Mali Bozo III; *Europe*: 1 Sardinian VI; 2 Italian VI IX; 1 German VI; 3 Basque VI, IX (2); *Asia:* 3 Japanese IV, V, VII; 2 Han Chinese VII, 1 Taiwan Atayal VII, 1 Taiwan Ami,

VII, 2 Cambodian VI, VII; *Pakistan*: 2 Hunza VI, IX; 2 Pathan VI, VII; 1 Brahui VIII; 1 Baloochi VI; 3 Sindhi III, VI, VIII; *Central Asia* 2 Arab IX; 1 Uzbek IX; 1 Kazak V; *MidEast*: 1 Druze VI; *Pacific*: 2 New Guinean V, VIII; 2 Bougainville Islanders VIII; 2 Australian VI, X: America: 1 Brazil Surui, 1 Brazil Karatina, 1 Columbian, 1 Mayan all X. An additional 1,009 chromosomes, representing 21 geographic regions, were genotyped by DHPLC for all markers other than those on the terminal branches of the phylogeny. The latter were genotyped only in individuals from the haplogroup to which those markers belonged. This hierarchic genotyping protocol was necessitated by the minute amounts of genomic DNA available for most samples.

[0099] **PCR**. The RepeatMasker2 program (http://ftp.genome.washington.edu) was used to identify human repeat DNA sequences. Primers were designed to amplify unique sequences and repeat elements other than LINE as confirmed by a negative female control, yielding amplicons 300-500 bp in length. All primers had a uniform annealing temperature, which allowed a single PCR protocol to be used. It comprised an initial denaturation at 95°C for 10 min to activate AmpliTaq Gold®, 14 cycles of denaturation at 94°C for 20s, primer annealing at 63-56°C using 0.5°C decrements, and extension at 72°C for 1 min, followed by 20 cycles at 94°C for 20 s, 56°C for 1 min, and 72°C for 1 min, and a final 5-min extension at 72°C. Each 50-µl PCR reaction contained 1 U of AmpliTaq Gold® polymerase, 10 mM Tris-HCl, pH 8.3, 50 mM KCl, 2.5 mM $MgCl_2$, 0.1 mM each of the four deoxyribonucleotide triphosphates, 0.2 µM each of forward/reverse primers, and 50 ng of genomic DNA. PCR yields were determined semi-quantitatively on ethidium bromide stained agarose gels.

[00100] **DHPLC analysis**. Unpurified PCR products were mixed at an equimolar ratio with a reference Y chromosome and subjected to a 3-minute 95°C denaturing step followed by gradual reannealing from 95 to 65°C over 30 min. Ten microliters of each mixture were loaded onto a DNASep™ column (Transgenomic, San Jose, CA), and the amplicons were eluted in 0.1 M triethylammonium acetate, pH 7, with a linear acetonitrile gradient at a flow rate of 0.9 ml/min^2. Under appropriate temperature conditions, which were optimized

by computer simulation (available at http://insertion.stanford.edu/melt.html), mismatches were recognized by the appearance of two or more peaks in the elution profiles.

[00101] **DNA sequencing**. Polymorphic and reference PCR samples were purified with QIAGEN (Valencia, CA) QIAquick spin columns. Both strands were sequenced to determine the location and chemical nature of any polymorphic sites, using the amplimers as sequencing primers and ABI Dye-terminator cycle sequencing reagents (PE Biosystems, Foster City, CA). Each cycle sequencing reaction contained 6 μl of purified PCR product, 4 μl dye terminator reaction mix, and 0.8 μl of primer (5 μM). Cycle sequencing was started at 94°C for 1 min, followed by 25 cycles of 96°C for 10s, 50°C for 2s, and 60°C for 4 min. The sequencing products were purified with Centrifex™ gel filtration cartridges (Edge Biosystems, Gaithersburg, MD) and analyzed on a PE Biosystems 373A sequencer.

[00102] **Statistical analysis**. The program CONTML in PHYLIP, version 3.57c, was used to construct a frequency based maximum likelihood network. The expected Luria-Delbrück/Lea-Coulson distribution of the number of mutants for each gene was fitted by maximum likelihood, treating each nucleotide of the screened sequence as analogous to a parallel, independent bacterial culture Luria, S. E. & Delbrück, *Genetics* 28:491-511 (1943); Lea, D. E. & Coulson, A. C. *Genetics* 49:264-285 (1949). The distributions under the expectation of constant population size were calculated according to Watterson, G. A. *Theor. Popul. Biol.* 7: 256-276 (1975). Mismatch distributions were calculated as described previously (Shen et al., *supra*). The NRY mutation rate per nucleotide per year (1.53×10^{-9}) was calculated on the basis of 597 nucleotide substitution differences between human and chimpanzee observed over 39,931 bp of non-coding sequence (Shen et al., *supra)*. The corresponding mutation rates for mtDNA (1.65×10^{-8}) and X chromosome (7.54×10^{-10}) were calculated on the basis of 581 and 58 nucleotide substitution differences, respectively, between human and chimpanzee observed over 6,176 bp of coding mtDNA (mictochondrial DNA) sequence

comprising the genes *ND1, ND2, COX1, COX2, ATP8, ATP6, COX3,* and *ND3*, and 7,853 bp of flanking non-coding sequence of the *DIAPH2* gene on Xq22.

[00103] **Accession numbers**. Most of the NRY sequence surveyed was derived from 5 cosmid sequences retrievable from Genbank using the accession numbers AC003031, AC003032, AC003094, AC003095, and AC003097. Six polymorphisms were affiliated with genomic regions for DFFRY (AC002531), one each for DBY (AC004474) and UTY1 (AC006376), 3 for SRY (NM003140), and 15 for random genomic STSs reported by Vollrath D, et al. *Science* 258:52-59 (1992).

[00104] The tree of Figure 2 is rooted with respect to non-human primate sequences. The 116 numbered compound haplotypes were constructed from 167 mutations (markers) of which 160 were discovered by DHPLC (Table 1). Seven haplotypes from the literature with less geographical heritage specificity were also analyzed in this study, including YAP (M1), DYS271 (M2), PN3 (M29), SRY 4064 (M40), TAT (M46), RPS4YC711T (M130), and SRY 2627 (M167), (the sequences for these markers are not shown in TABLE 1). Marker numbers indicated on the segments are discontinuous because of the removal of all but one polymorphism associated with tandem repeats and homopolymer tracts whose ancestral state is uncertain. Haplotypes are assorted into ten haplogroups (I – X) using principles commonly applied to haploid mtDNA phylogenies. Macaulay, V. et al. *Am. J. Hum. Genet*. 64: 232-249 (1999). Haplogroup I members, ancestral for M42, M94 and M139, also share the only homopolymer-associated marker M91. All haplogroup I individuals have an 8-T length variant, while 1,009 men in haplogroups II-X have 9 T's and in two cases 10 (not shown). Only one inconsistent haplogroup X individual had 8 T's (not shown). Haplogroups I and II, both of which are almost exclusively represented in Africa only, share the ancestral allele of M168. Haplogroup III is generally the most frequent one in Africa. Its frequency decreases with increasing distance from Africa, from 27% in the Mid-East to a few percent in Northern Europe and South and Central Asia. Haplogroup IV, related to the former through M1 and M145, is found mainly in Japan.

[00105] In a recent cladistic analysis of nine diallelic NRY polymorphisms, including M1, in 1,544 individuals, it was hypothesized that haplogroup III comprises descendents of a range expansion that brought Y-chromosomes back to Africa (M. F. Hammer et al. 15:427-441 (1998)). Haplogroups V and VIII are prevalent in New Guinea and Australia, but they are also found at varying though smaller frequencies throughout Asia. Haplogroups VI and IX are found mostly in Europe and the Indus Valley. They are not observed in East Asia, where haplogroup VII dominates, suggesting that this part of the world where agriculture developed independently resisted effectively subsequent gene flow Macaulay, V. et al *supra.* The distinction between Eurasians and East Asians was also observed with mtDNA Macaulay, V. et al., *supra.,* and autosomal genes (Diamond, J. *Guns, Germs, and Steel* (Norton & Co., New York, p. 99, 1999). Haplogroup X is common in the Americas, although its origin may have been in Central Asia where traces of it persist, as shown in Table 2:

TABLE 2.

Haplogroup	Exemplary Defining Mutation	Avg. no. of Mutations from Root to Individual Haplotypes	Total no. of Individuals	No. of Mutations per Haplogroup Minus Defining Mutation(s)	No. Haplotypes per Haplogroup
I	M91	6.1 ±0.95	52	20	8
II	M60	6.1±0.41	52	12	10
III	M96	10.4±0.24	218	27	21
IV	M124	10.5±0.56	9	7	4
V	M130	6.6±0.6	40	8	5
VI	M89 & absence of M9	7.4±0.25	163	25	23
VII	M175	9.5±0.35	137	18	15
VIII	M9 & Absence of M175 and M45	8.9±0.63	67	16	11
IX	M173	10.2±0.20	195	13	13
X	M74 & Absence of M173	9.2±0.1	129	6	6
Totals		8.59±0.20	1052	152	116

EXAMPLE 2

[00106] The root of the phylogeny was placed using sequence information generated from the three great ape species. The sequential succession of mutational events is unequivocal, except for those appearing in the same tree

segment (*e.g.*, M42, M94, M139). The phylogeny is composed of 116 haplotypes and their frequencies in 21 general populations are listed in Table 3. Forty-two haplotypes (36.2%) are represented by just one individual. Several haplotypes, however, display higher frequencies and/or geographic associations that reveal patterns of population affinities apparent from a maximum likelihood analysis (Figure 3) performed on the haplotype frequencies reported in Table 3. To facilitate presentation, the 116 haplotypes were grouped into 10 haplogroups as defined either by the presence or absence of mutations occupying strategic positions in the phylogeny. Haplogroups VI, VIII, and X, although polyphyletic, are distinguished by the criteria in Table 2.

[00107] Three mutually reinforcing mutations, M42, M94 and M139 (2 transversions and a 1-bp deletion) unequivocally distinguish haplogroup I which is represented today by a minority of Africans, mainly Sudanese, Ethiopians, and Khoisans (Table 2). All non-African, except a single Sardinian, and the majority of African males sampled, carry only the derived alleles at the three sites. This implies that modern extant human Y-chromosomes trace ancestry to Africa and that the descendents of the derived lineage left Africa and eventually replaced archaic human Y-chromosomes in Eurasia.

[00108] An important property of a phylogeny is the randomness of number of mutations per segment of the tree. Forty-one of the total 166 segments carry no mutation, while 98, 16, 8, 2, and 1 segment have 1, 2, 3, 4, and 8 mutations, respectively. The mean number of mutations per segment is 1.024 with a variance of 0.945. Applying the G-test for goodness of fit and Williams' correction to the observed G, the data do not fit a Poisson distribution (G_{adj}=34.98, df=3, P~10^{-7}). This is due to an excess of segments with one mutation, as expected in an exponentially growing population. Similar results were obtained recently for the separate analysis of 4 Y-chromosome genes. Further support that the human population has undergone a major expansion comes from the consistently negative values of Tajima's D (Lea, DE & Coulson, AC Genetics 49: 264-285 (1949)) for not only the Y-chromosome, but also for mitochondrial DNA, X-

chromosomal and autosomal genes. Interestingly, NRY shows evidence of significantly reduced variability to the other genetic systems (Shen et al., *supra*), confirming a similar comparison of a smaller number of polymorphisms on previously reported NRY sequences with eight X-linked (Hudson, R. et al, *Genetics* 116:153-159 (1987); Nachman, M. W. *Mol. Biol. Evol.* 15: 1744-1750 (1998) and 16 autosomal human genes. Possible explanations include positive selection on NRY Jaruzelska, J et al., *D. Mol. Biol. Evol.* 16:1633-1640 (1999) and a difference between male and female effective population sizes Wyckoff, G. J et al., *Nature* 403:304-309 (2000). Assuming expansion, the age of the most recent common ancestor (T_{MRCA}) was previously estimated at 59,000 years with a 95% probability interval of 40,000-140,000 years (Thomson, R. et al. *supra).*

[00109] This value is similar to an estimate of 46,000 to 91,000 years based on 8 Y chromosome microsatellites (Pritchard, J. K et al, *Mol. Biol. Evol.* 16:1791-1798 (1999) and, therefore, is considerably less than estimates of >100,000 years obtained previously (Hammer et al, *supra*). Of course, this assumes that selection or population structure have not had a major effect on NRY diversity, an assumption that may be wrong in light of our findings of significantly reduced variability on NRY. As the average number of mutations of all segments departing from the root is 8.60 (Table 3), and with a T_{MRCA} value of 59,000 years, the average time for adding a new mutation to the tree is 6,900 year. This puts the age of M168 that marks the expansion of anatomically modern humans out of Africa at approx. 44,000 years, in agreement with a previous estimate of 47,000 years with 95% probability intervals of 35,000 to 89,000 years using the program GENETREE (Thomson, R. et al. *Proc. Natl. Acad. Sci.* USA 97:7360-7365 (2000).

TABLE 3.

Haplotype Group	I								II										III																				IV
Haplotype #	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40
Sudan		17	1									5	1													2			1						7			2		
Ethiopia	6	5			1					3	1	4	1									3				15				16		2			20	6				
Mali								1	3		1			1					1	1		7						13	2								1	12		
Morocco																						2													1			1		
C. Africa											1	1			1	7	1	1			1	20					3													
Khoisan				11		5	1									11						7														4				
S. Africa				3								7										28	1	3	2		8			1						1				
Europe																																1								
Sardinia		1																														1			4					
Basque																																						1		
Mid-east																						2						1				1			2				1	
C. Asia + Siberia																																2	1		1					
Pakistan + India												2																			2			1						
Hunza																																								
Japan																																								1
China																																								
Taiwan																																								
Cambo + Laos																																								1
New Guinea																																								
Australia																																								
America																						2													1					
Total	6	23	1	14	1	5	1	1	3	3	3	19	2	1	1	18	1	1	1	1	1	71	1	3	2	17	12	14	2	19	2	7	1	1	36	11	1	16	1	2

Group	IV			V					VI																							VII									
Haplotype #	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81
Sudan																															4										
Ethiopia																															4										
Mali																																									
Morocco										1								3													14										
C. Africa																																									
Khoisan																																									
S. Africa																																									
Europe									1	1		8		1				2	1												9										
Sardinia										11						1															2										
Basque										2		1																			1										
Mid-east															1	2		2													8										
C. Asia + Siberia					10	16										2	1	12		4		1	1		2			1		1	17							6	9		2
Pakistan + India						1											4	3		3	2			1	1		1	4			7							1	1		
Hunza						1												1		3						1		2			1							1			
Japan	1	5	1			1	1	1																												2		2	1		6
China																																1			1	2	1	2	4	1	2
Taiwan																																	4			18					
Cambo + Laos						1																		2					1					1			1		1		
New Guinea				4																																					
Australia						3							1																												
America						1					1	1																			1										
Total	1	5	1	4	10	24	1	1	1	15	1	10	1	1	1	5	5	23	1	10	2	1	1	3	3	1	1	7	1	1	68	1	4	1	1	22	2	12	16	1	10

Group	VII					VIII											IX													X						
Haplotype#	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	98	100	101	102	103	104	105	106	107	108	109	110	111	112	113	114	115	116	Total
Sudan																																				40
Ethiopia																							1													88
Mali																							1													44
Morocco							1																5													28
C. Africa																																				37
Khoisan																																				39
S. Africa																																				53
Europe																	3	1					29				3									60
Sardinia																			2																	22
Basque																				2	7	5	26													45
Mid-east									2														2													24
C. Asia + Siberia		2			1	5			2	2	12	1											10			1	30	3	6	3		12	6			184
Pakistan + India							1	2	8	2													6		1		28			2		4				88
Hunza						2				3													3				11			2		7				38
Japan												1																								23
China		3	1	1																											1					20
Taiwan	5	46				1																														74
Cambo + Laos		1		1	6	1																					1									18
New Guinea						7							2	5	4	1																				23
Australia																							1	2												7
America						1	1																5							5				83	4	106
Total	5	52	1	2	7	17	3	2	12	7	12	2	2	5	4	1	3	1	2	2	7	5	89	2	1	1	73	3	6	12	1	23	6	83	4	1062

[00110] This concurs with recent archeological and mtDNA data, and is also consistent, though at a compressed time scale, with the weak Garden of Eden hypothesis. Under this hypothesis, a small subgroup of behaviorally modern humans left Africa and separated into several fairly isolated groups represented today by the major haplogroups III-X. Those groups remained small throughout the last glaciation before they underwent roughly simultaneous expansions in size as suggested by the star-like genealogy shown in Figure 1. In conclusion, the new levels of biallelic variation revealed here suggest a recent ancestry of the paternal lineages of our species from Africa and testify to the informativeness of the Y chromosome in deciphering the evolution of humankind.

[00111] The gene frequencies of New Guineans and Australian aborigines were grouped together because of the small sample size of the latter. Values at nodes indicate number of 1,000 bootstrap trees presenting cluster distal of node. Sudanese and Ethiopians are distinct from the other Africans and appear to be more associated with samples from the Mediterranean basin. This may reflect either repeated genetic contact between Arabia and East Africa during the last 5,000 to 6,000 years or a Middle Eastern origin with subsequent acquisition of Negroid genes on the way southwest with agricultural expansion. Native Americans are located between Eurasians and East Asian indicating common ancestry with both. This network is consistent with the first two principal components capturing 18% of the variation present in the 116 haplotypes.

EXAMPLE 3

[00112] A phylogenetic tree was deduced from NRY polymorphisms on the principle of maximum parsimony (Figure 3). Figure 3 shows the phylogenetic tree deduced from 304 polymorphisms including those presented in Examples 1 and 2 as well as other novel markers.

[00113] The contemporary global frequency distribution of the 10 Groups based on >1000 globally diverse samples genotyped using a hierarchical top down approach is illustrated in Figure 3. 171 haplotypes are identified in Fig.3 as well as their relationship with 309. However 4 markers are recurrent but define

distinctive haplotypes when considered in the context of the other markers. The 4 markers are M64.1 (M64.2),M108.1 (M108.2), M116.1 (M116.2) and 12f2.1 (12f2.2). For example M64.1 occurs on haplotype #80 in Group V and M64.2 on ht#159 in Group IX.

[00114] The relationship of the haplotypes to the ten haplogroups is also shown in Fig. 3. Each haplotype can be related to a specific geographical region within the haplotype group, allowing for very specific geographic association and ethnic identity of male individuals. Fig. 3 also shows which specific markers are important branching points for distinguishing between haplotype groups and also sub-haplotype groups such as haplotypes 10-13 of group II. This composite collection of 315 NRY variants (polymorphic markers) provides improved resolution of extant patri-lineages.

EXAMPLE 4

[00115] The methods of the invention can be utilized in the area of forensics to determine the ethnic affiliation of an individual.

[00116] The method involves, obtaining a nucleic acid sample from the individual and processing the sample sufficiently to conduct PCR amplification on the sample. The method exploits the hierarchical property of the Y chromosome gene tree that reveals the unequivocal sequential accumulation of DNA variation during the lineal life spans of these haplotypic molecules. Since Y chromosome haplotypes display a strong correlation with geography, such data provides insights into the affinity and diversification of populations. The sample is analyzed at polymorphic sites defining key internal nodes within the phylogeny. At least 11 primers sets, with each primer set recognizing at least one polymorphic region on the Y chromosome from a different haplotype group (Group I through Group X) are required to begin localizing a sample within the phylogeny. Additional haplotype resolution can be obtained by typing a subset of related markers. Each PCR reaction carried out on the sample, may include one or more primer sets/reaction vessel.

[00117] The PCR amplified products are analyzed by DHPLC (or any other suitable PCR product detection technique, such as DNA chips, direct sequencing, Taqman and the like) genotyping technology to define the haplotype which is then compared to a data base detailing the geographic association of the haplotype. The data base utilizes the markers identified in TABLE 1 and various combinations thereof which enables the identification of an individual to a particular haplotype group (Group 1 through Group X) as well as haplotype which are indicated in FIG.2 and FIG.4.

[00118] In certain instances, primer sets to the following markers are utilized to identify which haplotype group an individual originates from; Markers- M91, M60, M96, M174, (M216 or M316), M89, M9, M175, M45, M173. These markers identify the following haplotype groups; Group I = M91, Group II = M60, Group III = M96, Group IV = M174, Group V = M316, Group VI = M89 without M9, Group VII = M9 without M175 or M45, Group VIII = M9, Group IX = M173 and Group X is represented by marker M74 without M173. This approach can be expanded to increase criteria for inclusion/exclusion decisions.

[00119] TABLE 4 shows a two stage scheme of 30 markers, the haplotype groups they help define as well as geographical region associated with the haplotype group and the polymorphic markers which provides considerable power in facilitating localization any Y chromosome in the phylogeny. In cases where more than one marker is listed in TABLE 4, any one marker in the subset will provide comparable information.

TABLE 4

Markers analyzed Analysis #1	Assoc. Geographical region	Markers analyzed Analysis # 2	Assoc. Geographical region
M42, M94, M251, or M299 (Groups II-X)	Global	M215, M243, or M293 (Group III)	Africa, Med
M246 (Group I)	Africa	M2, M180 or M291 (Group III)	Sub Saharan Africa

M181 or M249 (Group II)	Africa	M191 (Group III)	Sub Saharan Africa
M168 or M294 (Groups III-X)	Global	M35 (Group III)	Africa, Med, S. Europe
M96 (Group III)	Africa, Med.	M217 (Group V)	E. Asia, India, N. America,
M174 (Group IV)	Asia, Japan	M201 (Group VI)	Med., S. Europe
M216 or M316 (Group V)	Asia, America, Polynesia, Melansia	M172 (Group VI)	Med., S. Europe
M89, M213 or M235 (Groups VI-X)	Global	M267 (Group VI)	Med., S. Europe
M9 (Groups VII-X)	Global	M170 or M258 (Group VI)	Europe
M175 or M214 (Group VII)	E. Asian	M52 or M69 (Group VI)	India
M45 or M74 (Groups IX-X)	Global	M122 (Group VII)	E. Asia
M173 or M207 (Group IX)	Europe, W. Asia	M119 (Group VII)	E. Asia
M269 (Group IX)	W. Europe	M268 (Group VII)	E. Asia
M242 (Group X)	C. Asia, India, Americas	M17 or M198 (Group IX)	E. Europe, W. Asia
M304 (Group VI)	Med.	M3 (Group X)	N.& S America

[00120] This example demonstrates that by using about 10% of the markers, one can localize any sample to a "neighborhood" or sub-haplotype group in the tree. These markers are useful in identifying a male for which no ethnic origin is

known. If it was known that the individual to be typed was for example, from Peking, then the assemblage of a more "Asian" group of markers would be more useful than those in TABLE 4.

[00121] The methods of the invention allow for the ability of Y markers to define (on a general geographic or population level) male ethnic affiliation.

[00122] While the present invention has been described with reference to the specific embodiments thereof, it should be understood by those skilled in the art that various changes may be made and equivalents may be substituted without departing from the true spirit and scope of the invention. In addition, many modifications may be made to adapt a particular situation, material, composition of matter, process, process step or steps, to the objective, spirit and scope of the present invention. All such modifications are intended to be within the scope of the claims appended hereto.

TABLE 1

M2 = DYS271 (209 bp) **A to G** at position 168
aggcactggtcagaatgaagTGAATGGCACACAGGACAAGTCCAGACCCAGGAAGGTCC
AGTAACATGGGAGAAGAACGGAAGGAGTTCTAAAATTCAGGGCTCCCTTGGG
CTCCCCTGTTTAAAAATGTAGGTTTTATTATTATATTTCATTGTTAACAAAAGT
CC**R**TGAGATCTGTGGAGGATAAAGggggagctgtattttccatt
For: 5'-3' = aggcactggtcagaatgaag
Rev 5'-3' = aatggaaaatacagctcccc

M3 = DYS199 (241 bp) **C to T** at position 181
taatcagtctcctcccagcaAGTGATATGCAACTGAGATTCCTTATGACACATCTGAACA
CTAGTGGATTTGCTTTGTAGTAGGAACAAGGTACATTCGCGGGATAAATGTG
GCCAAGTTTTATCTGCTGCCAGGGCTTTCAAATAGGTTGACCTGACAATGGGT
CACCTCTGGGACTGA**Y**AATTAGGAAGAGCTGGTACCTAAAATGAAAGATGCc
cttaaatttcagattcacaatttt
For: 5'-3' = taatcagtctcctcccagca
Rev 5'-3' = aaaattgtgaatctgaaatttaagg

M4 = DYS234 (273 bp) **A to G** at position 88
tcctaggttatgattacagagcgAGGATTATTATAATATTGGAATAAAGAATAATTGCTACA
AACTAATGATTAATGATATTCATAT**R**TAATCATATCTAAGATCTATATCTAGT
ATAACTATTCTTATTTTATATATTTTATTGTACTGGAACAGCTTGTGCCCTTGG
TCTCTTGCCTCGGCACCTGGGTGGCTTGCCATCCACAGAAGTGTTTTAACAGC
AAAAATTACTGTGAATTTTCTGCCCAAAAccttgtcatgtttacaagacgt
For: 5'-3' = tcctaggttatgattacagagcg
Rev 5'-3' = acgtcttgtaaacatgacaagg

M5 = DYS214a (322 bp) **C to T** at position 73
gggtttatactgacctgccaatgttAAAAGGGACCTAAATTCACTTTGGGGAAGTGGCCAGA
AAGGAAGAAG**Y**AGAAGGAGAAGAGTGCAAGAAACCTCCAGTTGTGGGGGTT
GAGCCTCCAGGATAAGAAAGAAAGAAATCTCCAGTAGGGGGGATTGAGCCT
AACACAAACCTTTGGTAATAGACAAGGCAAGACATTTCCAATAGGGGAGATT
GAGTGTCACCTCAAAACTATTAAGATGGGAAATACCCCAGGTAAGATAGAGG
GTAAAAAAGGATAAAGCTAGCAGCAATAACATTCcccctgaaagttcccaataa
For: 5'-3' = gggtttatactgacctgccaatgtt
Rev 5'-3' = ttattgggaactttcagggg
DYS214 complete. (656 bp) This fragment was converted into two STSs, a & b, containing M4 and M16 respectively. The two new STSs (a & b) omit an extra internal 68 bp region within the complete STS.
GggtttatactgacctgccaatgttAAAAGGGACCTAAATTCACTTTGGGGAAGTGGCCAGA
AAGGAAGAAG**C**AGAAGGAGAAGAGTGCAAGAAACCTCCAGTTGTGGGGGTT
GAGCCTCCAGGATAAGAAAGAAAGAAATCTCCAGTAGGGGGGATTGAGCCT

AACACAAACCTTTGGTAATAGACAAGGCAAGACATTTCCAATAGGGGAGATT
GAGTGTCACCTCAAAACTATTAAGATGGGAAATACCCCAGGTAAGATAGAGG
GTAAAAAAGGATAAAGCTAGCAGCAATAACATTCcccctgaaagttcccaataaTTTATG
CTAAAATATTGGAAAGACAACGAAAGGACTAAGCACAAGAGAAAGCAACAG
ATGATAAATATtgttatgtcatttgaacccagGAACCAATCTTCGAACCCTCAGTTTTCTGG
CCAAAGTTGGAGTCAAATGAGGATTGGATTTGTCAGCTTTTAATAGAACATA
TGATGACAAAACCTTCATCTCCCAGGAGGAGATAAATTATGCCCTATGTTGGT
GGCAAGGACCTGTCCTCCTTTACCCTCTAAAAACTGGAGGGAGAAAGTCAAA
GACTAACTCCTCTGAAAAAGATAAAGTCCCTATTCCTAgacagcccagcaacacacgg
For 3'-5' = gggtttatactgacctgccaatgtt
Rev 5'-3' = ccgtgtgttgctgggctgtc

M6 = DYS198 (218 bp) **T to C** at position 37
CactaccacatttctggttggCTTGTAGTTCTTTCT**Y**GGAAAAATATTATTCTAATTTCCTT
ATAGTATTAGCCATCAAAGTAGGGGAAGCAGATCAAATCTACCATAAGACCA
AGTCATAGGAAGAAGATCAAATTAAGATGCTAGGCAAAAGTCTCAGCACATA
TGGATTATGAGAAGCACATTCACACATCCAAActcaaagaatggactcagcg
For: 5'-3' = cactaccacatttctggttgg
Rev 5'-3' = cgctgagtccattctttgag

M7 = DYS253 (300 bp). **C to G** at position 236
ActgtgagcgagctgaaaatGCCTGATTTTCTCCCTTGGTTTAATGTAAAGGAAGGGATC
CAAAGGCTTAGGGAGATTGGGATGGTGGATTAGTCACTTTAGACCTACTCAT
TCCAATAGGGAGGGTCCAGAAGATGTACCCTTGACCAATGCCTTGCAAAATA
GATTCGTGAGGGCAGCACCTGCATCACCAAAGGGCATGTAATCATTCCTCTCT
GTATGTCAGATCTAACAASAAGAAGAACAGTAACTCAACTACAAAATTTAAA
CACAATGGAAAtaattggttcacaaggctgc
For: 5'-3' = actgtgagcgagctgaaaat
Rev 5'-3' = gcagccttgtgaaccaatta

M8 = DYS263 (267 bp). **G to T** at position 137
CccacccacttcagtatgaaTTTTGGGATCTGTTACCTATTTTTGATATAAAATCAACTG
CAAGTTTAGTGCCTCAGTATCACAAACACTGTATTTGCTCATATGTCTGTGAA
TCAATAACTTGGACTGGGTTCA**K**TTGGGCAGTTCTTCTATTGGTCTTGCCTGG
GGTCTTTAATGCAGCTTCCATTTTCTGGCAGCTTGATGAGACTGGATGGTCTA
AGGTACATTCATGAACACATCTGTTTGgtggacttgtctgtcagcct
For: 5'-3' = cccacccacttcagtatgaa
Rev 5'-3' = aggctgacagacaagtccac

M9 (340 bp) G10.35a **C to G** substitution at position 68
GcagcatataaaactttcaggACCCTGAAATACAGAACTGCAAAGAAACGGCCTAAGAT
GGTTGAATSCTCTTTATTTTTCTTTAATTTAGACATGTTCAAACGTTCAATGTC
TTACATACTTAGTTATGTAAGTAAGGTAGCGCTTACTTCATTATGCATTTCAA
TACTCAAAAAAAATTCCTTTGTGAAATGTTGAAATATTTTTCTAATCTGTTTC
ACGAGCTTCAAAAATGAGGAAAAAAGATTCAGTTTACATTTCAGCAAAATGC

CTCTTTTTAATCGGATTTATGTTTACTTAACATTTACAGTACATTTACgcttgagcaa
agttaggtttt
For: 5'-3' = gcagcatataaaactttcagg
Rev 5'-3' = aaaacctaactttgctcaagc

M10 = G10.10 (343bp). **T to C** at position 156
Gcattgctataagttacctgc**A**ATTTATAAAGTTGTGAAATAGTTCAAGACAATGAAGGG
AGAGACTCTCTGGTAACTACAGAGTATGAGCTCATCATTGCTTAGTTTCCACA
AGAGGTATCTCTGAATTTTTTTGTTTATTCCCAATGATCTTA**Y**AGCACTTGTA
AAGTTTTTACATTAGTTACAAAATGCAATTTGAAGTGAAAGAAACAGAAATA
CAAAATATTAGTTTCTCTTTTTCTCCTACATTCCTACATGGATTTGTAGAAGAG
CTGACCTTTACTTATAAAATAAATCAGCAAATGAGTGTCTTTTCTAGAATGggg
tgacccaatttttatta
For 5'-3 = gcattgctataagttacctgc
Rev 5'-3' = taataaaaattgggtcaccc

M11 = G10.37 (222 p) **A to G** at position 44.
Tctctctgtctgtctctccctcc**C**TCTCTCCTTGTATTCTAAC**R**GAAAGGTTTAGAACTTGCA
TAATTGGGAAAGAAGCTGTTGCCTGAACTTACTGGGGGATTCAGCATTGTCA
TTTTGGACATGTCACTTATCCTCAGTATTTGCTTCCCCCAGGAGAGAGCTGTA
ATAAAAAAGCATTGCAATTTAATACATAAgctcagtaagttcttgtttatgctc
For: 5'-3' = tctctctgtctgtctctccctcc
Rev 5'-3' = gagcataaacaagaacttactgagc

M12=DYS260a (309 bp) **G to T** at position 286
Actaaaacaccattagaaacaaagg**A**CTTAAACTAGGAATTAATTATTTCTCTTTCTCTTTC
CATGGCCAACAAACATTGAAAAAAAATTGCCATCTTTTTTTTTATTTGTTTGTT
AGAGATGGGGATCTCACTCTGTTTCTTAGATTGTAGTGCCATGGCACAATAAT
GGCTCACTGCAGCCTCAAACTCCTGGGCTCAAGTGATCACCCCCATACAGAC
TCCCGAGTAGCTGGGAACACAGGCACATGCCACCACCCCTAGCTAATTTTTT
ATTATTTGTAGA**K**ATGggggtcactatgttgctcag
For: 5'-3' = actaaaacaccattagaaacaaagg
Rev 5'-3' = ctgagcaacatagtgacccc

M13 = G10.06 (233 bp) **G to C** at position 157
Tcctaacctggtggtctttc**A**TTGTTTTACAAAGGTGATTTAGTTTTGGGAAGGACTATTC
TCCTTTAAACTATAGACTAAATTTTTCTCAAAGTTAGGTTAGTTTATGCCCAG
GAATGAACAAGGGCAGTAGGTAGGTTAAGGGCAAGACGGTTA**S**ATCAGTTCT
CTGTTACTGTTATAATTTTCTCATTGTTATATTTTTTGCAAATGTGgttggataaaatca
tggctca
For: 5'-3' = tcctaacctggtggtctttc
Rev 5'-3' = tgagccatgattttatccaac

M14 = G10.07 (287 bp) **T to C** at position 180
Agacggttagatcagttctctg**T**TACTGTTATAATTTTCTCATTGTTATATTTTTTGCAAAT
GTGGTTGGATAAAATCATGGCTCATACAAATATACAAAAAATACATATTAAA

ATTTTATTTAACATAAAACATTAAAATTTATTTAATAAATTATAAATGAAAAA
ATCAGTAACATG**Y**TATAAGCAGTTTAAAAAAGTTAATGAAGCTCAGTTTTAA
CATGAAGTATAGGAATGGTGAAATTATATAAATGAAATTTGTAAATggtgtcaatgt
gcttttatcta
For: 5'-3' = agacggttagatcagttctctg
Rev 5'-3' = tagataaaagcacattgacacc

M15 = G10.16 (295 bp = ancestral state); derived allele = **9 bp insertion** (304 bp) after position 109; Note that there are also two T to G changes immediately before the 9 bp insertion.
AcaaatcctgaacaatcgcCATCACCTATTTGGTGGACGCATAGGCCTGGTCTCTGATCT
GGTCGCATGTCCAGAGGGTCTGCTAACCCACTGCACCTAGGGAGACA**TTGTA**
CAGAGACATTGTACCACCTTTTCTCTACTcttcccagactcaacacatttGATTGTATATGC
GCATGAGGTAGAAATATAAGATGAAGCAGGGACAGAGTCAACAAGCCAGAA
CTAGATGCTTCTACCTGGACAGAAGACCTAGAATTCTTTTTTGGATCCTAAAT
TCACCAggaaattttaaccacatgca
For: 5'-3' = acaaatcctgaacaatcgc
Rev 5'-3' = tgcatgtggttaaaatttcc
M15 polymorphic region in more detail
mutant sequence = GACA **TT GTACAGAGA** CA
ancestral sequence = GACA GG * * * * * * * * * CA

M16 = DYS214b (266 bp) **C to A**
TgttatgtcatttgaacccagGAACCAATCTTCGAAC**M**CTCAGTTTTCTGGCCAAAGTTG
GAGTCAAATGAGGATTGGATTTGTCAGCTTTTAATAGAACATATGATGACAA
AACCTTCATCTCCCAGGAGGAGATAAATTATGCCCTATGTTGGTGGCAAGGA
CCTGTCCTCCTTTACCCTCTAAAAACTGGAGGGAGAAAGTCAAAGACTAACT
CCTCTGAAAAAGATAAAGTCCCTATTCCTAgacagcccagcaacacacgg
For: 5'-3' = tgttatgtcatttgaacccag
Rev 5'-3' = ccgtgtgttgctgggctgt

M17 = G10.47a (333 bp) **-1bp deletion** (4G's to 3G's) at position 68
CtggtcataacactggaaatcAGATTCTGTCTACTCACCAGAGTTTGTGGTTGCTGGTTGT
TACGGG**G**TTTTTTTAAGTGAATTTTGGGGTTTGTTAAGTGGCCAAACTATTTT
TGTGAAGACTGTTGTATGTGGGTTTCAGATGTCTCTACATCAGTTTGTGGTCA
GCTAGTGAGTTAAATTTTATGAAAAGCCTGGAGAAACAAGAATAGCAGTAAA
AACTTCCAGTCTTTGTAGATTGGGTGTCTTCAGTGCTTAGCTGGGCAATTTAA
AACTTACCTTAAGTAGTACAGTTGGCCCTTTGTGTCTGTgagtttcacatttgtaggttca
For: 5'-3' = ctggtcataacactggaaatc
Rev 5'-3' = tgaacctacaaatgtgaaact

M18 = G10.47b (333 bp = ancestral size) **+2 bp (extra AA) insertion** after position 62
CtggtcataacactggaaatcAGATTCTGTCTACTCACCAGAGTTTGTGGTTGCTGGTTGT
T**AA**ACGGG**G**TTTTTTTAAGTGAATTTTGGGGTTTGTTAAGTGGCCAAACTATT
TTTGTGAAGACTGTTGTATGTGGGTTTCAGATGTCTCTACATCAGTTTGTGGT
CAGCTAGTGAGTTAAATTTTATGAAAAGCCTGGAGAAACAAGAATAGCAGTA

AAAACTTCCAGTCTTTGTAGATTGGGTGTCTTCAGTGCTTAGCTGGGCAATTT
AAAACTTACCTTAAGTAGTACAGTTGGCCCTTTGTGTCTGTgagtttcacatttgtaggttc
a
For: 5'-3' = ctggtcataacactggaaatc
Rev 5'-3' = tgaacctacaaatgtgaaactc

M19 = G10.47c (333 bp) **T to A** at position at 131
ctggtcataacactggaaatcAGATTCTGTCTACTCACCAGAGTTTGTGGTTGCTGGTTGT
TACGGG**G**TTTTTTAAGTGAATTTTGGGGTTTGTTAAGTGGCCAAACTATTTT
TGTGAAGACTGTTGTA**W**GTGGGTTTCAGATGTCTCTACATCAGTTTGTGGTC
AGCTAGTGAGTTAAATTTTATGAAAAGCCTGGAGAAACAAGAATAGCAGTAA
AAACTTCCAGTCTTTGTAGATTGGGTGTCTTCAGTGCTTAGCTGGGCAATTTA
AAACTTACCTTAAGTAGTACAGTTGGCCCTTTGTGTCTGTgagtttcacatttgtaggttca
For: 5'-3' = ctggtcataacactggaaatc
Rev 5'-3' = tgaacctacaaatgtgaaactc

M20 = G10.48. (413 bp) **A to G** at position 118
GattgggtgtcttcagtgctTAGCTGGGCAATTTAAAACTTACCTTAAGTAGTACAGTTGG
CCCTTTGTGTCTGTGAGTTTCACATTTGTAGGTTCAACCAACTGTGGATTGAA
AAT**R**TTTGAAAAATTAAAAATAGATGGTTGCATTTGCACTGAACATGTAGAC
TTTTTTTTCTTGTAATTTCTCTTAAACCATACAGCATAACAACTCTTTACATAG
CATGTACATTGTATTAGGTATTCTGAGTACTCTAAAGTATACGGGAGGATGTG
TGTAGGTTATGTGCAAATACTATAACATTATATGTAAGGGATTTGAAAATTCT
GGGATTTTGGTATTTGCAGGTGGTGTGGGATGGGGGTCTGCCTGGAACCAAG
GAATGCCCCAAAGGAGgatggtgccttgttgtgtg
For: 5'-3' = gattgggtgtcttcagtgct
Rev 5'-3' = cacacaacaaggcaccat

M21 = G10.43 (415 bp) **A to T** at position 357
CttttatttctgactacagggCCCTCTTTTGCATTGTTTTGTAGGTCAGATTTATTAGTAGT
ATGTTCTTTCAGCTTTTGTGTATCTGGGAATATTTCAGTTTCTCCTTTATTTTG
AAGGATAGTCTTTGAGTTTTTCCTACTTAACAGATCCTGGAGCTTCTTGGATG
TGTAAATTAATGATTTTCATCAAATGTGAAGTTGTTTTCGGCTATTCTGCAGA
TATCCTTTACCACCCCTTTGCTGCCTCTTCCTATTGTGGGTAATAGGCATGTCT
CTGTATGTTGGAGAGAATCAAAGGTCTTTTAAGCCCTTGATTTTTATTTATCTT
TTGTTTTTTGTTCCTCAGACTGTAT**W**GTTTCAGTTGACTTAGCTTCCAGTTTGT
TGATTCTTCTGcctgctcaaatctgctgtt
For: 5'-3' = cttttatttctgactacaggg
Rev 5'-3' = aacagcagatttgagcagg

M22 = DYS273 (327 bp) **A to G** at position 129.
AgaagggtctgaaagcaggtTCGTGATTTCACCCTTTACAGTTTAATACAAGGGATTTTA
CATACAGACATATAAGCTGATAGTCCTGGTTTCCCTATTTGTTTTAAGGTGCC
ATTCCTGGTGGCTCT**R**CCTCCTTCCCCCAGTGCCCATATGGGCCCTTAGTCTG
CTGTAGGCATGCTCAGGCAAGCCCTTGAGCAAATTCCCTTAATCTGCACGAA

ACATGGGCTGGAGATTCAGTGGGACCCTTTCTTTAGTGTCTGCCTAATGCAAG
CTGGCTAACTCCTTTCAAAAGTTTTGTCTTGCTGATgaagcctccaggtagtaggc
For: 5'-3' = agaagggtctgaaagcaggt
Rev 5'-3' = gcctactacctggaggctt

M23 = G10.57a (327 bp) **A to G** at position 159
TctctaacttctgtgagccacTCTAGCAAATTAATTGAACCAAAGGAGGAGGTTAAGGAC
AGCATAGTTTACAAAATGAGCCCTGTTTCTGACATCTGAAGTGGGGGCAGTC
TAGTGGGCCTGACCTCTTAACTTGTAGAAACATTCTTTCTTTCTAG**R**TGACTA
GTGACCAGAATTAAATTGAATCCTAGGCCACCCATTTATTGTCTTCTGCAGAA
TTGGC**G**AGAATGGAGAGGAATCCTCACCTATC**G**GTGACCAGAGATGAAATAT
TCTGAATTGAGAGTTTAAAAGAGCACACTTAGAagagatttagagtttagtttttcc
For: 5'-3' = tctctaacttctgtgagccac
Rev 5'-3' = ggaaaaactaaactctaaatctct

M24 (tetranucleotide TAAA motif) = SRY 8299c. Internal primer regions for SRY4064 which contain M40 and M41.
Acagcacattagctggtatgac**A**GGGGAGATGTGATTAATTGACCTACTGATAAGACTCA
TTTCAGTAAATGCCACACAAGAATgtataataggctgggtgctgTGGGTCACACCTGTAA
TCCCAGCCCTTCGAGAGGTCAAGGCGA**G**CGGATCACAGGGTGGAAGAGATT
GAGACCATCCTGGCCAACATGGTGAAACT**G**GGTCTCTACTAAAAATACAAAA
AATTAGCTGGGCGTGGTGACATGTGCCTGTAATCCCAGTTACTCGGGAGGCT
GAGGCAGaagaatcatttgaactcatgAGGCAGAGGTTGCAGTAAGCTGAGATTGCGCCG
CTGCACCCCAGCCTGGCAACAGAGCGAGACTTTGTCTCAAAAA**A**AAA**W**AAA**T**
AAATAAATAAATAAATAAACAATAATAAAAAAAGCGTAATAGCTAGCCTATC
CTACCCTATATTCTAAAATTCAAAAGTAATGGTTTTTGTTATGAAATCTcgtaagt
cttgccataaagaga
For: 5'-3' = acagcacattagctggtatgac
Rev 5'-3' = tctctttatggcaagacttacg

M25 = B9.008b. (340 bp) **G to C** substitution. Position 121
Aaagcgagagattcaatccag**G**ATGACAGAATGCGTTCACCTTTAAAGGGATTAAAAGA
AGTATAATACAGTCTGTATTATTAGATCACCCAGAGACACACAAAACAAGAA
CCGTGAATT**S**AATTAGTGGTATACTAATAGAGTGGTTTTACCTGAAATATTTA
CACATCAATCCTACTGAATTCTTACAACAAATGATTTAGATTAGCTATTGTAT
TCACCAGTTGAAAGAACAGAAAATATTGAGGGAGATAACTTGTGTCAGTGCA
ACTTAATCAGATTTAGGACACAAAAGC**A**ACTACATAATGAAAAAGAGAgctggt
gacttaacttgctaaaa
For: 5'-3' = aaagcgagagattcaatccag
Rev 5'-3' = ttttagcaagttaagtcaccagc

M26 = B9.005 (321 bp) **G to A** at position 68
CcagtggtaaagttttattacaatttTTTTAAACCAAGATTCAATTTTTTTCTGAATTAGAATT
ATC**R**CAGAGAACACTGAATGGCCTATGAAATTCAATTTTTGCTGCAGATTTC
GTCATGTTTCTTAATGAACATATAACTAACTTCTAATCACAAGATAAATTCTT
GCCTATGTGCAAAAACTTAGTGCTGCATCCTTGTGTATGGTTTTAAAAAGTGT

CAAAACTGGCCCCTCATGTCAAATACAGCCCCAATTAGGGGAGGCAACCTAA
GAAAGGTGTACAACTGTCCTGACATTggattgcctgcttactgtgaa
For: 5'-3' = ccagtggtaaagttttattacaatttt
Rev 5'-3' = ttcacagtaagcaggcaatcc

M27 = G10.65. (526 bp). **C to G** at position 398.
CggaagtcaaagttatagttactggAAATACAAACTGTGGCAGTAGAAAACCCTAGGCACA
AGGGAAGTAAAATATTAACCACTCCAGGCTGGAGTGCAGTGGCGCAATCTGG
GCTCACAGCAAGCTCTGCCTCCTGGGTTCACACCATTCTCCTGCCTCAGGCTC
CCGAGTAGCAGGGAGTACAGGCACCCGCCACCAGGCCTGGCTAGTTTTTTTT
GTATTTTTTAGTAGAGATGGGGTTTTACTGTGTTAGCCAGTATGGCCTCGATT
TCCTGACCTCGTGATCCGCCCACGTCAGCCTCCTAAAGTGTGGGGATTACAG
GAGTGAGCCACCATGCCCAGCTGAAACAATAGTTCTTCACAATGGCATCTAC
CACTATGTCCACATTTGCACCT**S**TGTCCTGAACCTCGATTCCTATAGGTTGAT
GTGTTGAGAACCAGACAATACGAAATAGAAGACAAATCATGAGCTTACAGA
ACCTGAAACTTTTTACACTGGGCAGtgtggtagacagaacagcagtg
For: 5'-3' = cggaagtcaaagttatagttactgg
Rev 5'-3' = cactgctgttctgtctaccaca

M28= G10.33n (332 bp). **T to G** at position 277.
GcttacttgggacacaggctAGTTCTCTCCTGAAGCTATTGAGCAGTATGTGTTGAGGTG
CGCTACGCCAGTTGAGGTGAAGCTGTTACACAGTATGAAAGCCGGGCTTTGT
AGCTGCAGCTGCGCATTGCACCCCCAGCTACGCAGTCTCCTTTCCTTCTCAGT
CACAGGACCGGATGGCAAGTGGCCGCAGCCAGTCGGTGAGACCGACTGAGC
TCTGGGGCTTCAGTTCTTGACGCTACCTACATGGCTACATCTCCAGCCAAGGA
TGAGAGG**K**GATGCCAGAGGACCTCGATCTAAATTGGGCAccattatcgtatgacaacttct
ct
For: 5'-3' = gcttacttgggacacaggct
Rev 5'-3' = agagaagttgtcatacgataatgg

M30 = G10.66 a (486 bp) **G to A** at position 132.
GaaccagacaatacgaaatagaagACAAATCATGAGCTTACAGAACCTGAAACTTTTTACA
CTGGGCAGT**G**TGGTAGACAGAACAGCAGTGGCTGCCCAAAGATGATCATGTT
TTAAGTCCTGACATCTGT**R**AATTATCATATTGGGAAAAGGTGTTATTGTAGAT
GTTGTTTAAAGTTAGGATTTTGAGAGAGGAAAATTATGTAGGGTTATCTGGCT
GTGCCCAGTGAAATCACAAGAATCTTTATAAATGAAAAAAGAAAGCAGAAG
AATCAGAACCAGAGACACGGCATTATGCATAGGACTGGACTTGTCATTACTA
GTTTTAAAGGTAGAGGAAGCAGAGATCTAAGAAATGCAGGCAGCCTCTAACT
AATGTTAACAAATCTCATTTTCTAATATTGTAAGCCTGTGGAAGAGGCTAGGG
CACAGATGCTCCCATAGAGTCTCCAGAAGGAACCTAAggtaatgagataagccgctaaa
For: 5'-3' = gaaccagacaatacgaaatagaag
Rev 5'-3' = tttagcggcttatctcattacc

M31 = G10.66 b (486 bp) **G to C** at position 71.
GaaccagacaatacgaaatagaagACAAATCATGAGCTTACAGAACCTGAAACTTTTTACA
CTGGGCAGT**S**TGGTAGACAGAACAGCAGTGGCTGCCCAAAGATGATCATGTT

TTAAGTCCTGACATCTGT**G**AATTATCATATTGGGAAAAGGTGTTATTGTAGAT
GTTGTTTAAAGTTAGGATTTTGAGAGAGGAAAATTATGTAGGGTTATCTGGCT
GTGCCCAGTGAAATCACAAGAATCTTTATAAATGAAAAAAGAAAGCAGAAG
AATCAGAACCAGAGACACGGCATTATGCATAGGACTGGACTTGTCATTACTA
GTTTTAAAGGTAGAGGAAGCAGAGATCTAAGAAATGCAGGCAGCCTCTAACT
AATGTTAACAAATCTCATTTTCTAATATTGTAAGCCTGTGGAAGAGGCTAGGG
CACAGATGCTCCCATAGAGTCTCCAGAAGGAACCTAAggtaatgagataagccgctaaa
For: 5'-3' = gaaccagacaatacgaaatagaag
Rev 5'-3' = tttagcggcttatctcattacc

M32 = G10.68a (370 bp) **T to C** at position 166.
TtgaaaaaatacagtggaacAAAGATCCTCTGTATCTCTGCTCCTAAGATAGCAGAGACA
GCatactggcttctgttcaattttcCTTTGATTACACAACTTCATTGGCTACGGTGTTTAATAT
GACCGTCATAGGCTGAGACAAGATCTGTTCAGTTTATCTCA**Y**AAGTTACTAG
TTA**A**ATCTCAGACATATTATACTTTTGTAACTGAGTGACTCCCATTGTAAGGA
TAACTACTTCAATGTGCGTATAAATGAGTCAGTTGTCTCTCTTGGGGGCTTCA
ACAAATAAGCAAAGATAACCTCATTGTGGAGAGCACTTCACATTTGTTTTTAG
GGTTACATAGTCTActctgtatccttaaacacttg
For: 5'-3' = ttgaaaaaatacagtggaac
NewF 5'-3' = atactggcttctgttcaattttc
Rev 5'-3' = caagtgtttaaggatacaga

M33 = G10.68b (370 bp) **A to C** at position 180.
TtgaaaaaatacagtggaacAAAGATCCTCTGTATCTCTGCTCCTAAGATAGCAGAGACA
GCatactggcttctgttcaattttcCTTTGATTACACAACTTCATTGGCTACGGTGTTTAATAT
GACCGTCATAGGCTGAGACAAGATCTGTTCAGTTTATCTCA**T**AAGTTACTAGT
TA**M**ATCTCAGACATATTATACTTTTGTAACTGAGTGACTCCCATTGTAAGGA
TAACTACTTCAATGTGCGTATAAATGAGTCAGTTGTCTCTCTTGGGGGCTTCA
ACAAATAAGCAAAGATAACCTCATTGTGGAGAGCACTTCACATTTGTTTTTAG
GGTTACATAGTCTActctgtatccttaaacacttg
For: 5'-3' = ttgaaaaaatacagtggaac
New F 5'-3' = atactggcttctgttcaattttc
Rev 5'-3' = caagtgtttaaggatacaga

M34 = G10.69 (372 bp) **G to T** at position 131
CacttcacatttgtttttaggGTTACATAGTCTACTCTGTATCCTTAAACACTTGAAGATCT
GTTATAACTACATCTGAGATAGTAGTCACAGTGTTTTCTCATGTTAATGCCTG
GCTTCCACCCAGGAG**K**CACATGTGGTGTGTCTGCAAATAAAGTGTTTATGAT
TATTGGGGTCCCCCAAGCTGGACCTGTATCCATGTTCAAGTGGCCACAGGGTT
ACTTGCTTTAGCATGGCTCCTTGGCTGGCTGTTAAGTGAATAATTAAACTGAG
TCTTTTTTGCAGGAGCTAACTGAGACCAATCAATCAGTCAATTTTCCCTTTCT
GTGTGTAACACAAGCTGGATGTCCctggaatgactaaataatgact
For 3'-5' = cacttcacatttgtttttagg
Rev 5'-3' = agtcattatttagtcattccag

M35 = G10.72a (514 bp) **G to C** at position 168

TaagcctaaagagcagtcagagTAGAATGCTGAATTTTCAGAAGTTTTATATTAACATAA
TCATTCATCTTTTTTGTCCTGATAATTACTCAGGAGGAAACTGAGAGGGCATG
GTCCCTTTCTATGGATAGCAATACTCAGTGTCCCAATTTTCCTTTGGGACACT
G**S**GACACAGGCAGAGACTCCGAAAGTCTGCATGGATTAGTTGTTCATTCACC
ACAGCTCCTTAGTGTGCCAGGAGAACTATATATGGCCTTTGGTTTCATTCAGG
GACAGGGAAACTTGAACCCATGCCTATTCATTCTCATTAAAGTAGCAGAAGT
CATGTTAGAGACAGTATTGCTGCATTCAGTACTCCTGCCTTTAACGCTTCTGA
CGCTTCCTGAAAGCAGCCCCAGCTCTCCATATGGCAAAACAAAGGCAACCTT
ATGCAAAGCCTTCTCAGGGAACCCTCAGAAAGGTTTAAACTTAGGTTCACAG
TTTTTAGAGAATAAtgtcctcattgctccctctg
For: 5'-3' = taagcctaaagagcagtcagag
Rev 5'-3' = cagagggagcaatgaggaca

M36 = G10. 82a (436 bp) **T to G** at position 74
AgatcatcccaaaacaatcataaCTTGTTTAAATTGTTCATAGCAAAAGTTACATATTATA
AAGAGTTATGAG**K**GTCTTAGGCAGTGAATAGTAACTGAATATCCTTTTATAG
TTGTCCTTCACTAGCAGGAAGCCTTATTCCCTGCCCTTTTACATATCTTAACTT
AGAATGTTACTGTCTAAATAGTGGTTAGGCAAGAGTAGTTCTTAAACGTGCA
GTAATTATCTTGCACTACATTTAAGGGCTAAATAGCTAGTAGTGGTGCTTGAT
AATTGAAGAAATTTGTACAGCTGGAGGAAGTACCTGCTAAATTTTCAAAAGT
TACCTGAATTTAATAGGTAAATCTGTTTTTAATTAGAGCTATATCATTTTACTC
TGAATGTCTTAACATA**G**AAGTTTACATAAAATTTAcagattggattgatttcagcctt
For: 5'-3' = agatcatcccaaaacaatcataa
Rev 5'-3' = aaggctgaaatcaatccaatctg

M37 = G10.STS 84 (422 bp) **C to T** at position 203. This STS also contains M61 at position 101 which is defined in G10.83.
CagattggattgatttcagccttCTTCTGGTACTTTTTAAAATCTTATTAATCATTAGGAAAA
GAAGTTTTATTATTGATGCAAGCCCTAAACACTCTTTCGACTCCAGAGGAGAA
GCTGGCAGCTCTCTGTAAGAAATATGCTGATCTTGTGAGTATTTATTTAATGG
AGCAAGGAACACAGAAAATAAAATCTATGTGTG**Y**TTGATAAGATTTTTAAAT
ATTATTTTGATGTAACTTTAAATGTAAAATGATATTTTATCTCAAAATTGAAA
ACAATCTCCTTTCTTTAGTACTTATGATTGGTGTGTGTGACTTCATCTTATGAA
ATGATGTATAGAACATAATAATACTTTTTTAAATGTGAAATAAATTTCCTAAA
ACTTAATATGCTAGATCAgcagtttttttttttgtatgct
For: 5'-3' = cagattggattgatttcagcctt
Rev 5'-3' = agcatacaaaaaaaaaaactgc

M38 = G10.73a (337 bp) **T to G** at position 146
CagtttttagagaataatgtcctCATTGCTCCCTCTGGCACTAGCAGTTTGTACCAGGAGAT
CTGTTGGCTACTGTTACCCTAGGGTATGGCAATGGTATGTAGGCAATGAAAA
ATCTTACAGTACTTATTATGGAAAACCAACT**K**TTTTATTCAGTAAGCATTCCC
CTGTGTTGTAAGGTTTTTAAAAGATTGTGGAAGTATGAAAAAGTTTATTATGA
CAGATGTGCCAGCTC**C**AGCTGTTTTGTGGAGAGTGACCCTTGGATTTTCGTAT
GCCCCCATTATATGATGATACCTTGTAATGATTTAATTTTAGcatctgcttttctttttctttaa
For: 5'-3' = cagtttttagagaataatgtcct

Rev 5'-3' = ttaaagaaaagaaaagcagatg

M39 = G10.73a (337 bp) **-1 bp (-C) deletion** at position 236
CagtttttagagaataatgtcctCATTGCTCCCTCTGGCACTAGCAGTTTGTACCAGGAGAT
CTGTTGGCTACTGTTACCCTAGGGTATGGCAATGGTATGTAGGCAATGAAAA
ATCTTACAGTACTTATTATGGAAAACCAACT**T**TTTTATTCAGTAAGCATTCCC
CTGTGTTGTAAGGTTTTTAAAAGATTGTGGAAGTATGAAAAAGTTTATTATGA
CAGATGTGCCAGCTC**C**AGCTGTTTTGTGGAGAGTGACCCTTGGATTTTCGTAT
GCCCCCATTATATGATGATACCTTGTAATGATTTAATTTTAGcatctgcttttctttttctttaa
For: 5'-3' = cagtttttagagaataatgtcct
Rev 5'-3' = ttaaagaaaagaaaagcagatg

M41 = SRY 4064b (218 bp) **G to T** at position 117. Site is located within SRY 8299 509 bp STS.
GtataataggctgggtgctgTGGGTCACACCTGTAATCCCAGCCCTTCGAGAGGTCAAGG
C**A**AGCGGATCACAGGGTGGAAGAGATTGAGACCATCCTGGCCAACATGGTG
AAACT**K**GGTCTCTACTAAAAATACAAAAAATTAGCTGGGCGTGGTGACATGT
GCCTGTAATCCCAGTTACTCGGGAGGCTGAGGCAGaagaatcatttgaactcatg
For: 5'-3' = gtataataggctgggtgctg
Rev 5'-3' = catgagttcaaatgattctt

M42 = B9.008a (340 bp) **A to T** substitution at position 297
Aaagcgagagattcaatccag**G**ATGACAGAATGCGTTCACCTTTAAAGGGATTAAAAGA
AGTATAATACAGTCTGTATTATTAGATCACCCAGAGACACACAAAACAAGAA
CCGTGAATT**G**AATTAGTGGTATACTAATAGAGTGGTTTTACCTGAAATATTTA
CACATCAATCCTACTGAATTCTTACAACAAATGATTTAGATTAGCTATTGTAT
TCACCAGTTGAAAGAACAGAAAATATTGAGGGAGATAACTTGTGTCAGTGCA
ACTTAATCAGATTTAGGACACAAAAGC**W**ACTACATAATGAAAAAGAGAgctgg
tgacttaacttgctaaaa
For: 5'-3' = aaagcgagagattcaatccag
Rev 5'-3' = ttttagcaagttaagtcaccagc

M43 = DYS260b (309 bp) **A to G** at position 77
Actaaaacaccattagaaacaaagg**A**CTTAAACTAGGAATTAATTATTTCTCTTTCTCTT**T**TC
CATGGCCAACAAAC**R**TTGAAAAAAAAATTGCCATCTTTTTTTTTATTTGTTTGT
TAGAGATGGGGATCTCACTCTGTTTCTTAGATTGTAGTGCCATGGCACAATAA
TGGCTCACTGCAGCCTCAAACTCCTGGGCTCAAGTGATCACCCCCATACAGA
CTCCCGAGTAGCTGGGAACACAGGCACATGCCACCACCCCTAGCTAATTTTTT
ATTATTTGTAGA**G**ATGggggtcactatgttgctcag
For: 5'-3' = actaaaacaccattagaaacaaagg
Rev 5'-3' = ctgagcaacatagtgacccc

M44 = G10.87 (389 bp) **G to C** at position 263
Ctggcaccttctgatattttgag**A**AGCAGGAATCCCTGAGCATAAATGTAAATAGCTTAGA
ACTGTCCAAAAGCAAAGACAGCAGAAAATAAAATTGTTGCTTGCTATGTTCA
GGAAAGGAATGCTTCCATTGGATATGGAAGCCAGTCTCAATTGTTACATCAG

CCTGAGGAAACTCATGCGAGAAATGCCAGAAAAAGAAGACAGCAACAAAGA
AGATAAAAGAAAGACTGACAAAAGCATTGAATTTCTGGTAGAAAAA**S**CAGT
GTACTAGAAGGTTAGGAGATTTCCTAGCTGTCAGCCATGAAAGGGTTGGGGA
AGAAAGAGCAATTTGGTTGCATACTGTAGCATGGTCATCTAGGGTGgtcctcaaac
acatagaaatcaca
For: 5'-3' = ctggcaccttctgatattttgag
Rev 5'-3' = tgtgatttctatgtgtttgaggac

M45= B9. 12(352 bp) **G to A** substitution at position 109
GctggcaagacacttctgagCATCGGGGTGTGGACTTTACGAACCAACCTTTTAACAGTA
ACTCTAGGAGAGAGGATATCAAAAATTGGCAGTGAAAAATTATAGATA**R**GC
AAAAAGCTCCTTCTGAGGTCCAGGCCAGGAGATAGTAGGATTTAAGAAACAA
ACAAACAAAAAC**A**ACCACAAATGACCTTTGGTGCCACTGTCACAACTGTTGC
TCATCAGAGTAGGAGAGTTGTAGCAAAGGCATTAAAGAAGGACAAGCAGCT
GAAGAGCCTGAATCCTTGTGTTGTAAGCTATTTTGGTTTCCTTTCAAGAAAGG
GCTGTGGTCTGTggaaggtgtcaggaacatatt
For: 5'-3' = gctggcaagacacttctgag
Rev 5'-3' = aatatgttcctgacaccttcc

M47 = G10. 82b (436 bp) **G to A** at position 395
AgatcatcccaaaacaatcataaCTTGTTTAAATTGTTCATAGCAAAAGTTACATATTATA
AAGAGTTATGAG**T**GTCTTAGGCAGTGAATAGTAACTGAATATCCTTTTATAGT
TGTCCTTCACTAGCAGGAAGCCTTATTCCCTGCCCTTTTACATATCTTAACTTA
GAATGTTACTGTCTAAATAGTGGTTAGGCAAGAGTAGTTCTTAAACGTGCAG
TAATTATCTTGCACTACATTTAAGGGCTAAATAGCTAGTAGTGGTGCTTGATA
ATTGAAGAAATTTGTACAGCTGGAGGAAGTACCTGCTAAATTTTCAAAAGTT
ACCTGAATTTAATAGGTAAATCTGTTTTTAATTAGAGCTATATCATTTTACTCT
GAATGTCTTAACATA**R**AAGTTTACATAAAATTTAcagattggattgatttcagcctt
For 5'-3'= agatcatcccaaaacaatcataa
Rev 5'-3' = aaggctgaaatcaatccaatctg

M48 = G10. 79n (240 bp). **A to G** at position 160
AaacaatatgtatgctaattttgctTAAAAGATTATACACTGAAATTTAGAGAGGATATAATG
TTATCTGTAGTGTAGAAAGAGTTAAATAAGACTGATTTTTAGAATTTGTTTTA
TCCCTTCCACTCTTAGCTTGACAATTAGGATTAAGAATATGAT**R**TGTCAAATT
TCATGACTGAAATCTGAAATGCCTTAATAGTTGCCCTCAGTGTTTcatccttatactaa
catttacattga
For: 5'-3' = aaacaatatgtatgctaattttgct
Rev 5'-3' = tcaatgtaaatgttagtataaggatg

M49 = B9.15new a (354 bp) **T to C** at position 229
CggcaacagtgaggacagtAGCTCCAGGTCTGGGCGGAAGGTGGTGCGGTGAAAGGTG
CAGGGACAGACTGGGTTAGAGGCCACTCTTGGTCTTATCCTCCATGGCCACA
ACAGAGGTGACAAATACATGGGTCACTCAGTTATGTTTAGCCAACAGCCTAC
CCAAACCACACC**T**GTCTTACCAGAGCCCTTTCCTGGAGCCATGTTCTCAGGAC
TGGTCACACTGTC**Y**CCATTCTCCAGCAGCCCTTGGACCTATCGGAAAAAAAG

AATGGGTAACAATAATTGAGCTGATGAACCAGGTCCTATCTTTCCTCCCACAA
CTCCAAAACTTGGgagcctctatctcctgaagca
For 5'-3'= cggcaacagtgaggacagt
Rev 5'-3' = tgcttcaggagatagaggctc

M50 = B9.15new b (354 bp) **T to C** at position 175
CggcaacagtgaggacagtAGCTCCAGGTCTGGGCGGAAGGTGGTGCGGTGAAAGGTG
CAGGGACAGACTGGGTTAGAGGCCACTCTTGGTCTTATCCTCCATGGCCACA
ACAGAGGTGACAAATACATGGGTCACTCAGTTATGTTTAGCCAACAGCCTAC
CCAAACCACACC**Y**GTCTTACCAGAGCCCTTTCCTGGAGCCATGTTCTCAGGA
CTGGTCACACTGTC**T**CCATTCTCCAGCAGCCCTTGGACCTATCGGAAAAAAA
GAATGGGTAACAATAATTGAGCTGATGAACCAGGTCCTATCTTTCCTCCCACA
ACTCCAAAACTTGGgagcctctatctcctgaagca
For: 5'-3' = cggcaacagtgaggacagt
Rev 5'-3' = tgcttcaggagatagaggctc

M51 = B9.16 (339 bp) **G to A** at position 33
GagcctctatctcctgaagcAGAGTAGACACA**R**GCTTCCAACAGGGATCAGAGTTTAGG
GATCTGGATAGGTATAGAATGGAGCAAAGGGACTAGGCCAAAGGAGATTGA
AAACTGGGGAACAGGGACAAGACTGGAGCTACAAGAAGGACAGGGGCTAGA
AGACAGAAATATGAGGACAATGGCTGGCCTGGAAAGCTCACCTTAGAAATAT
TGTTGCCACTGCCTTCTCTGATAGGGTCACAGGCAGTGGCTGAAGTGTAGACT
GAGGCCTCCTCTGGTCTGGGTTTGGCCTGTAGCTGTTGGCGAAGCTCAGCCAG
Ctgtcgcaacagagcagtca
For: 5'-3' = gagcctctatctcctgaagc
Rev 5'-3' = tgactgctctgttgcgaca

M52 = G10.88 (534 bp) **A to C** at position 477
Actgtagcatggtcatctagggtg**G**TCCTCAAACACATAGAAATCACACAAGAATTGTCAA
ATTGAAGATTTGGATTTAGTAGATCTGAAAACGCACTTTGTAAAATTGGCCAC
AGTAGAGGTGGAAGTGACTGAAATACTGCATTATTTATTTATTTAATTAATTT
ATTTTAGTCAGAGTCTTGCACTGTCGCTAAGGCTGGTATACCATGGTTCAGTC
ACAGTTCACTACAGTCTTGAACTCCTAGGCTCAAACAATTCTCCTGTATCGGC
CTCCTGAGTACCTGGCACTACAGACATGCACAAGCATGCATGGCTAATTTTA
AAAAAATTTTTGTAGAAATGGAGTCATGAACTCCTGGGCTCAAGTGATCCTC
CCACCTCAACTTCCCAGAGTGTTGAGTGAGATTACAGTTATGAGCCACCATCC
CTGGCCAATAAAGGTGTTTTTAATACCTATAAGAATATTGCCTGCA**M**GGATG
TTTGATAGGTTTCTTGATATTTCATTCTctctcttgaaatgtttgcttcgtc
For: 5'-3' = actgtagcatggtcatctagggtg
Rev 5'-3' = gacgaagcaaacatttcaagagag

M53 in tree **(tetranucleotide TAAA motif)** = SRY 8299d. Internal primer regions for SRY4064 which contain M40 and M41.
Acagcacattagctggtatgac**A**GGGGAGATGTGATTAATTGACCTACTGATAAGACTCA
TTTCAGTAAATGCCACACAAGAATgtataataggctgggtgctgTGGGTCACACCTGTAA
TCCCAGCCCTTCGAGAGGTCAAGGCGA**G**CGGATCACAGGGTGGAAGAGATT

GAGACCATCCTGGCCAACATGGTGAAACT**G**GGTCTCTACTAAAAATACAAAA
AATTAGCTGGGCGTGGTGACATGTGCCTGTAATCCCAGTTACTCGGGAGGCT
GAGGCAGaagaatcatttgaactcatgAGGCAGAGGTTGCAGTAAGCTGAGATTGCGCCG
CTGCACCCCAGCCTGGCAACAGAGCGAGACTTTGTCTCAAAAA**A**AAA**T**AAA**W**
AAATAAATAAATAAATAAACAATAATAAAAAAAGCGTAATAGCTAGCCTATC
CTACCCTATATTCTAAAATTCAAAAGTAATGGTTTTTGTTATGAAATCTcgtaagt
cttgccataaagaga
For: 5'-3' = acagcacattagctggtatgac
Rev 5'-3' = tctctttatggcaagacttacg

M54 = B9.17 (360 bp) **G to A** at position 164
Cctcctctggtctgggttt**G**GCCTGTAGCTGTTGGCGAAGCTCAGCCAGCTGTCGCAACA
GAGCAGTCACATCTTCAGAGGCCAGAGCCTTTCTGGCACGGTCTTGCCAGCC
AATGGCCCTCTCTGTGAGACACTGAAGGGCCTCACCCTCAGGCAGCCGCAC**R**
GGCAGCCTCTGCAGGGCAACCAGCAAGGCTAGGATTGTCTCTAGGCGTGGCC
GTCGTGAGCGCATACACAGTGGACACAGGAATTTTGTGTCCCATTCCCACCA
GGCTAGCAGTGGAGATGAAGTGAGACTGGGCTTTGGAGAGGTGAGGAGATG
GGGCACTGACACACACTGCCCatggaaccagtcctgacaca
For: 5'-3' = cctcctctggtctgggttt
Rev 5'-3' = tgtgtcaggactggttccat

M55 = B9.28 (382 bp) **T to C** at position 228
Cgtaggcgtttgacagcag**T**TAATAGAGACTACAGATATCAAAGTCAGAGAGTCCAGCT
TCCTGAGAAAACGTTAACAGTATTAATCTGCTACCACTATGGCTACTAATACC
ATGCCACCACGGTACTACCTGGCTAGTACCATTCCACAGAAGAACAGAAATA
AATACAAATAGGTGGGGCAAGAGAAAAGAAACATGTGAAAAGGCCCCTGGA
TGGTTTAAGTTA**Y**ATTTTCATCAGTCATCCAGTTAAGAGTTAAAGAATGAGG
AAGAGATGTAAAAACAGCCATTAGGATTCAGAAGTAGTAGCTTTCACAGTGA
GACAAAACATCTATTAAGCCAGAAACTGAAGTACAAATGCAATgggaggattacgaa
gaaagg
For: 5'-3' = cgtaggcgtttgacagcag
Rev 5'-3' = cctttcttcgtaatcctccc

M56 = B9.29 (399 bp) **A to T** at position 39
CcagaaactgaagtacaaatgcAATGGGAGGATTACGA**W**GAAAGGAGGGCTAAGTGAT
GATAAGTATGGTCAGAATAATAAATTTATTCTAGACAAGAAATGAGAGTTCA
TTATGTCAGAAGCAAAATAGTACTACAGGATGACAACTTCTGAGATTTACTCT
TTGGTTCCAACTGCCTACAAGACAAAGAAAACTGAAGAGGCCAGGAAGTTAA
ATGCATGAGGAAAACTTGAGGCAGATTAAAATGGAAATGCAGGGCATGTTAT
TTGGGTATCATGGGTTCAATCTGGAAAAGCCTTATTTCTCCTGAACCACAGTA
GGGAAAGGAGTTATCCAGAAAAGTGAAATTTATTCTAAAATTTTAAGTTTCC
ATGTTTTaaagagaggcagcaatgaga
For: 5'-3' = ccagaaactgaagtacaaatgc
Rev 5'-3' = tctcattgctgcctctcttt

M57 = G10.85n (326 bp ancestral); **+1 bp insertion** (327 bp = Derived). Extra A inserted at positon 133
Attgggaggaagtggtttctg**T**ATTTAAAATTTTCCGAAGGAATTCTGCAGATTCAAGCTC
TAACCATTCTTGATTAAAATTGTGAGTTAGATAAGATTGTTTAGTAAAATTGT
ACTATGGCTCAGGAAATA**A**TTTATTTAATATCTACTGTATGCCAAGCATTGTT
CTTTTTTCCATCTTCCAGGGAAATTCACCTCTTCTATAGAAGAGTTTGTTTTGA
ACTATACGATTTGAAACAAAATTCTTTTTTTGGAGACTATGGAAACATTCTCA
ACAGGGAAACCCTACTAGACTTTGTAAAgcaaataatggaaaagatacagaac
For: 5'-3' = attgggaggaagtggtttctg
Rev 5'-3' = gttctgtatcttttccattatttgc

M58 = G10.57b (327 bp) **G to A** at position 224
Tctctaacttctgtgagccac**T**CTAGCAAATTAATTGAACCAAAGGAGGAGGTTAAGGAC
AGCATAGTTTACAAAATGAGCCCTGTTTCTGACATCTGAAGTGGGGGCAGTC
TAGTGGGCCTGACCTCTTAACTTGTAGAAACATTCTTTCTTTCTAG**A**TGACTA
GTGACCAGAATTAAATTGAATCCTAGGCCACCCATTTATTGTCTTCTGCAGAA
TTGGC**R**AGAATGGAGAGGAATCCTCACCTATC**G**GTGACCAGAGATGAAATA
TTCTGAATTGAGAGTTTAAAAGAGCACACTTAGAagagatttagagtttagtttttcc
For: 5'-3' = tctctaacttctgtgagccac
Rev 5'-3' = ggaaaaactaaactctaaatctct

M59 = B9.15new c (354 bp) **A to C** at position 279
Cggcaacagtgaggacagt**A**GCTCCAGGTCTGGGCGGAAGGTGGTGCGGTGAAAGGTG
CAGGGACAGACTGGGTTAGAGGCCACTCTTGGTCTTATCCTCCATGGCCACA
ACAGAGGTGACAAATACATGGGTCACTCAGTTATGTTTAGCCAACAGCCTAC
CCAAACCACACC**T**GTCTTACCAGAGCCCTTTCCTGGAGCCATGTTCTCAGGAC
TGGTCACACTGTC**T**CCATTCTCCAGCAGCCCTTGGACCTATCGGAAAAAAAG
AATGGGTAACA**M**TAATTGAGCTGATGAACCAGGTCCTATCTTTCCTCCCACA
ACTCCAAAACTTGGgagcctctatctcctgaagca
For: 5'-3' = cggcaacagtgaggacagt
Rev 5'-3' = tgcttcaggagatagaggctc

M60 = B9.34 (388 bp ancestral); **+1 bp insertion** (389 bp = DERIVED). Extra T inserted after positon 242
Gcactggcgttcatcatct**G**GGAGCAGCTCAAAAGCCTCTCGCTCAGCCTCCGTGACGCC
CTGGGGGTGTTCAACCCACATATACTGTAAAGACTAGGAGTAGGGTTGTGGA
CACCCCACCTCAGCCAACACTGAGCCCTGATGTGGACTCAACCTTGTAAGGA
AAGCTGTAGAGAAATTGGAAGAAAAAATATAAACACATACAGACTCTGTCTT
TACATTTCAAAATGCATGACTTAAAG**T**ATCAGGCACACAGTGGTTACTCAAT
GTTGGTCTGTGTCTCTGTAACGTAATATATGTGACTAAATCCCTAAGCTCTGC
TCTTGACCACCCACCTTCTCCAAAAGGGCCTTTCGTAGACGTCGCTcctcctgaacca
taatgaacat
For: 5'-3' = gcactggcgttcatcatct
Rev 5'-3' = atgttcattatggttcaggagg

M61 = G10. 83new a (190 bp) **C to T** at position 98.

AttggattgatttcagccttcTTCTGGTACTTTTTAAAATCTTATTAATCATTAGGAAAAGA
AGTTTTATTATTGATGCAAGCCCTAAACACTCTTTYGACTCCAGAGGAGAAG
CTGGCAGCTCTCTGTAAGAAATATGCTGATCTTGTGAGTATTTATTTAATGGA
gcaaggaacacagaaaataaaat
For: 5'-3' = attggattgatttcagccttc
Rev 5'-3' = attttattttctgtgttccttgc

M62=DYS260c (309 bp) **T to C** at position 60
ActaaaacaccattagaaacaaaggACTTAAACTAGGAATTAATTATTTCTCTTTCTCTYTC
CATGGCCAACAAACATTGAAAAAAAATTGCCATCTTTTTTTTTATTTGTTTGTT
AGAGATGGGGATCTCACTCTGTTTCTTAGATTGTAGTGCCATGGCACAATAAT
GGCTCACTGCAGCCTCAAACTCCTGGGCTCAAGTGATCACCCCCATACAGAC
TCCCGAGTAGCTGGGAACACAGGCACATGCCACCACCCCTAGCTAATTTTTT
ATTATTTGTAGAGATGggggtcactatgttgctcag
For: 5'-3' = actaaaacaccattagaaacaaagg
Rev 5'-3' = ctgagcaacatagtgacccc

M63 = B9. 22 (308 bp) **G to A** at position 43
CtcttcccttggttcctattcTGACACGCTCAGGTACCTCAARGAATCCTCCAACTTCCCAC
CTTCACTTTCTAGCACAACCCAACCGAGTAAAAACTATAAAGTATATCTATCT
CTCTTCTAACTGCTGGCCTGACGCAGTAAAGCAGAAATACTGATCCTCACTTG
GATCTCATCCACATCAGCAATCCAAGCTTGTGCCTTAGTCAGAGCTTCTTTGA
GAGCCTGGATGTTAGGCAGGTGAACAGGGATGTTTTCTGTCTCACGAATTAT
GGCTTCCAATGTGGCTggtggatgcttctgcctaa
For: 5'-3' = ctcttcccttggttcctattc
Rev 5'-3' = ttaggcagaagcatccacc

M64 = B9.t23 (325 bp) **A to G** at position 279 **RECURRENT**
TatagaccctgactactcaagagaaAAGTCCAATCCAAAGAAAAAATACAAAAGAAAACA
AAATCACATCAGGCCACAAACCAGTTTAAGGGCCCTCACCACATGGTTGGCT
CCAGACTGAAACATTTCATAGGGGTAAATAATGCGTTCGTAATGTGATCGTA
GCAGGGAGCCAATGTTTTTGCCTGGTGGGTAGTGGAGACGCTGGGCAACTCG
AGCCCACCGACGATCCTTGCAGATGGCTTCATAGCCACCTTCCTCAATCACAA
TCTGAAAGTRTAAGAAACAATATGGATGAACTGTGAacagactggaaagggctacc
For: 5'-3' = tatagaccctgactactcaagagaa
Rev 5'-3' = ggtagccctttccagtctgt

M65 = B9.t26 (436 bp) **A to T** at position 152
TtctgatgccagcttgttcgGGTCAGAAAAGTTAAATGAGAAATTTGGTGCTAAGGGTTT
CTGGTCATGAGTGTAAATAACGCCTCGCCAAGTGGTAAACTGCCCCAACGTT
CAAACCAAAGGCTACCCATTCCCAAATTTTGTTTCAAAGWCTTACCGCGGGT
GGGCGGATTTTGCAGATGCCAGACTTCTCTGCTATGGGCCTTATTTTCGCAAT
GTAGCCAAGCGGGTCTTGGAATTCAGCCCAGCTAGGCTCAAAAACCGGGCAC
TCCGGTGGCGGCAGGAACTCGTCACACCCCGGTTCCATGTCGGGCCTTAATG
CTAAGCTGTAAAATAAGAATCACATTGTCTTTAATGACGCGCTGGTTCCTCCT
ACTAAAAGGCCTATGAAAATTTCATTTTCTTGAGAATTTcaaggttactttaatcccgtagc

For: 5'-3' = ttctgatgccagcttgttcg
Rev 5'-3' = gctacgggattaaagtaaccttg

M66 = B9.41 (415 bp) **A to C** at position 135
CtgtgtaacaccatcaagtgcACCCATATATGCAGAATGGGAATTTCGTAAGAAAAGAGA
AGGAAAAAGGCAGAACAGTTGAAGCAAAAATGGTTAAACAATTTCCAAATTT
GTGGAAAGCCCTGAAAGTCTAC**M**ACCAAGAAGCTCAGTGCACTCCAACTAG
ATAAACTCCAGGAGACACAACATAGTCGAACCAACAAAAGGTAAGACACCA
AGATGGAGTTTGAAAGCAGTATGACAGACATGATTCTTCGCATATAATGGAT
GCTTAATAGAATTATCAATAGATTTCTCATTAGAAATAACGGAGGCCAGAAG
CCAGTTGGATGACACGTTAAAAGTCATGCAATGGGAAAAAAAATTAAATAAA
TTGACAGAGAATTAAAAATTGTggaagtatgtctccagaagatgt
For: 5'-3' = ctgtgtaacaccatcaagtgc
Rev 5'-3' = acatcttctggagacatacttcc

M67 old = B9.36new a (409 bp) **A to T** at position 377
CcatattctttatactttctacctgcAGGCCCACTGCATGCTCACTCACCCAGTCAGCAGTACA
AAAGTTGACAGCTTCAGCAAAATTGTAGCCTTGGTTAAAACCACTGTGGTAA
GCACGAGGAAAAGTGATGACAAACTCCCCTGCACACTGGTTTGTGCGGACAA
CCTAAAAAGGAGAAAAAAGCAGAAAGAGGTGTGGGTCAGAACTAATGGGCC
AGATGTGAACTCAAAGATGTCTCTAGATGCTGTAACAGATGTAGGAAG**A**GTG
GAAAGGCTCTATCTTCAAGTACGTGTCCTAAAAGAAAAATGAGATTGTGAAT
TTAAAA**G**TGGTATTCATAGAAAAGTACTCAAAATATGTGTAATTCAAAAAAC
A**W**ATATAGAGGGgtccacgaacaagtgaaaagac
For: 5'-3' = ccatattctttatactttctacctgc
Rev 5'-3' = gtcttttcacttgttcgtggac

M67 revised B9.36new a (386 bp) STS **A to T** at position 327
ccagtcagcagtacaaaagttgACAGCTTCAGCAAAATTGTAGCCTTGGTTAAAACCACTG
TGGTAAGCACGAGGAAAAGTGATGACAAACTCCCCTGCACACTGGTTTGTGC
GGACAACCTAAAAAGGAGAAAAAAGCAGAAAGAGGTGTGGGTCAGAACTAA
TGGGCCAGATGTGAACTCAAAGATGTCTCTAGATGCTGTAACAGATGTAGGA
AG**A**GTGGAAAGGCTCTATCTTCAAGTACGTGTCCTAAAAGAAAAATGAGATTG
TGAATTTAAAA**G**TGGTATTCATAGAAAAGTACTCAAAATATGTGTAATTCAA
AAAACA**W**ATATAGAGGGGTCCACGAACAAGTGAAAAGACTCTttgcttctataatcaa
agaaatgc
newFor 5'-3' = ccagtcagcagtacaaaagttg
newRev 5'-3' = gcatttctttgattatagaagcaa

M68 old = B9.36new b (409 bp) **A to G** at position 268
CcatattctttatactttctacctgcAGGCCCACTGCATGCTCACTCACCCAGTCAGCAGTACA
AAAGTTGACAGCTTCAGCAAAATTGTAGCCTTGGTTAAAACCACTGTGGTAA
GCACGAGGAAAAGTGATGACAAACTCCCCTGCACACTGGTTTGTGCGGACAA
CCTAAAAAGGAGAAAAAAGCAGAAAGAGGTGTGGGTCAGAACTAATGGGCC
AGATGTGAACTCAAAGATGTCTCTAGATGCTGTAACAGATGTAGGAAG**R**GTG
GAAAGGCTCTATCTTCAAGTACGTGTCCTAAAAGAAAAATGAGATTGTGAAT

TTAAAA**G**TGGTATTCATAGAAAAGTACTCAAAATATGTGTAATTCAAAAAAC
AA**A**TATAGAGGGgtccacgaacaagtgaaaagac
For: 5'-3' = ccatattctttatactttctacctgc
Rev 5'-3' = gtcttttcacttgttcgtggac

M68 revised B9.36new b (386 bp) STS **A to G** at position 219
ccagtcagcagtacaaaagttgACAGCTTCAGCAAAATTGTAGCCTTGGTTAAAACCACTG
TGGTAAGCACGAGGAAAAGTGATGACAAACTCCCCTGCACACTGGTTTGTGC
GGACAACCTAAAAAGGAGAAAAAAGCAGAAAGAGGTGTGGGTCAGAACTAA
TGGGCCAGATGTGAACTCAAAGATGTCTCTAGATGCTGTAACAGATGTAGGA
AG**R**GTGGAAAGGCTCTATCTTCAAGTACGTGTCCTAAAAGAAAATGAGATTG
TGAATTTAAAA**G**TGGTATTCATAGAAAAGTACTCAAAATATGTGTAATTCAA
AAAACAA**A**TATAGAGGGGTCCACGAACAAGTGAAAAGACTCTttgcttctataatcaaa
gaaatgc
newFor 5'-3' = ccagtcagcagtacaaaagttg
newRev 5'-3' = gcatttctttgattatagaagcaa

M69 = B9.62a (257 bp) **T to C** at position 222
GgttatcatagcccactatactttgGACTCATGTCTCCATGAGAACTAAGACTACCACAACA
GAATCCCTATAGTCCAGCCCTCAGATCACATACATGTACAGGCATGTTGAAG
TAGTCGGACTTGAAGGAATCAGCCATTTCACCAAAACTCTGCAAACTGTACT
CCTGGGTAGCCTGTTCAAATCCAAAAGCTTCAGGAGGCTGTTTACACTCCTGA
AA**Y**AAAATATATTTCagcaagacaaagggaataaagat
For: 5'-3' = ggttatcatagcccactatactttg
Rev 5'-3' = atctttattccctttgtcttgct

M70 = B9.62b (257 bp) **A to C** at position 45
GgttatcatagcccactatactttgGACTCATGTCTCCATGAGA**M**CTAAGACTACCACAACA
GAATCCCTATAGTCCAGCCCTCAGATCACATACATGTACAGGCATGTTGAAG
TAGTCGGACTTGAAGGAATCAGCCATTTCACCAAAACTCTGCAAACTGTACT
CCTGGGTAGCCTGTTCAAATCCAAAAGCTTCAGGAGGCTGTTTACACTCCTGA
AA**T**AAAATATATTTCagcaagacaaagggaataaagat
For: 5'-3' = ggttatcatagcccactatactttg
Rev 5'-3' = atctttattccctttgtcttgct

M71 = B9.63b (328 bp) **C to T** at position 197
TtgaattatagtcccttgcctcTGGTTCAGTCAAGTCTCTATCATTCTAGAGTTAGTGTGTT
CAATCGTTCTTGTATAGTAGCTCACTGATAGCTTAATCAAAACCTAACACAAA
TATTAACTTATAAAAGGGCAGAAACTACCTTCCCAAAACCCA**G**AAGGGGAGA
TTACAGAAAATCACCAACCAAAAATAAAG**Y**ATCTGTGACAGACAGATCTTAC
CGCCAAGATACATTTTGGGCACCTCCAGATGCCTCTGGGGATTTCAGGAAGG
GGTGGTAACAAGCAGAAGATGTGGTAATTGTCATCAcagccatcacagaaaagaagc
For: 5'-3' = ttgaattatagtcccttgcctc
Rev 5'-3' = gcttcttttctgtgatggctg

M72 = B9.63a (328) **A to G** at position 157

TtgaattatagtcccttgcctcTGGTTCAGTCAAGTCTCTATCATTCTAGAGTTAGTGTGTT
CAATCGTTCTTGTATAGTAGCTCACTGATAGCTTAATCAAAACCTAACACAAA
TATTAACTTATAAAAGGGCAGAAACTACCTTCCCAAAACCC**R**GAAGGGGAG
ATTACAGAAAATCACCAACCAAAAATAAAG**C**ATCTGTGACAGACAGATCTTA
CCGCCAAGATACATTTTGGGCACCTCCAGATGCCTCTGGGGATTTCAGGAAG
GGGTGGTAACAAGCAGAAGATGTGGTAATTGTCATCAcagccatcacagaaaagaagc
For: 5'-3' = ttgaattatagtcccttgcctc
Rev 5'-3' = gcttcttttctgtgatggctg

M73 = B9.47a (361 bp ancestral & 359 bp derived) **-2bp deletion,**
(-GT) at position 260
cagaataataggagaatttttggtCAAATAAAAGGCCATATTATATTTCTTTTGATAAAAGT
ATCATGTGTTCAGTATGTTTTATTATTTGAAATAATTAACATGACAGGAATAT
ATTTGAAAAAAATTCCAAAAAAAGCTAAATATACAAACTAAGAAAATTATAT
GATTATACTTATCTGCAGTATTGTAAAACAATAGTTCCAAAAACTTCTGAATT
ACAAGTTTAATACATACAACTTCAATTTTC**A**ACTACATT**GT**GGTTAGACGTTC
AGAGGAATCACAAAGGACCTCAACATGCTAGATAAGAAAATGTATTTTTTAA
ATGTTTTGGCTCAgctgcttagaaaataaggaaaat
For: 5'-3' = cagaataataggagaatttttggt
Rev 5'-3' = attttccttattttctaagcagc

M74 = B9.50a (385 bp) **G to A** at position 195.
AtgctataataactaggtgttgaagATAAAATCAGTTTAATTTAAATAAGAGGATAAAAGAA
GTATGAGCAGAAAAAGGTTTTCAATATTAACTAGGAAAGTCTGAAAAATAAT
CAGAAATTCTAAAGATAAAAACATAACATTAAAAATTATAAACTAAGTTGTT
TAATAGATTAGGTATTTTAAAAACTGGT**R**CATTTTTAAGTTGCTTTAAGTAAG
TTACTTAAAAGACAACAGCAGCAAAA**G**AATTAAAAAAAAATGAAAGGTGAA
GAAACACATACAAGAGAACCTTAGAACAGTAAGGTTCTAGCTAACAGGAGA
AATAAATTACAGACTGTAAAAGTTGATGACCAAGAATTTTttcagaagtggtaaaagctg
aatt
For: 5'-3' = atgctataataactaggtgttgaag
Rev 5'-3' = aattcagcttttaccacttctgaa

M75 = B9.51 (355 bp) **G to A** at position 296
GctaacaggagaaataaattacagacTGTAAAAGTTGATGACCAAGAATTTTTCAGAAGTGG
TAAAAGCTGAATTCTCAAGTTTGAGAATTCCTATCTATTCCCAGAAATATTAA
GTAAAAAGTCACATTCCACACATCAAGAAAACTTGCAAGACACTAAAAGAG
ATATTATAGCAGTCAAATAGAAAAAGCAAAATAGACTACTACAAATTAATGT
AAGATTCAGAATTGACTTGTCAAAAGCCAAAACAGATTTCTAATGTACTGTG
AAAAGACAATTATCAAACCACATCC**R**TATATATACAGAGAAATACCTTTATA
AGAATAAAAATtcacaaatgcctctgttcaata
For: 5'-3' = gctaacaggagaaataaattacagac
Rev 5'-3' = tattgaacagaggcatttgtga

M76 = G10.100a (493 bp) **T to G** at position 339

TagaagtagcagattgggagaggACATGTGTTCAAGTTGTACTACTTGTATGTCTTGTTTA
GATATTACAGTCTTTTTCTTTTATCAGAAAATAATTGAATAATGATAAAATCA
GTTGCAGATTAAGACAGATTATCTGTTGCAGTCTTCTCAAAACTTAATTTAAG
TACATTATTTTCAGCTAGCATTTCTTCCTTCACATAGAACCTCCATGTGTGGA
GGGATTTCCTAATGAGTCTATTGTATGTACAATAGCACTTAATGACATAGCTT
TTAAATAATAACAGGATTTTACCAAATGTTTAATATGTGCCAGGCATCAAGC
ACC**T**TACACAGTT**K**AATTATTGCATAGATTTGGACAGCAACTCTGCAAGTTA
GGTATGGTCATGAACCTTTGCAGATAAGGAAACTGTGTTTCACAAGGAGAAG
AAATTGTCCTGGATCATACAATAAGCTAGGATTTGCTCCAgaccatttttttcattttatcagg
For: 5'-3' = tagaagtagcagattgggagagg
Rev 5'-3' = cctgataaaatgaaaaaaatggtc

M77 = G10.105 (371 bp) **C to T** at position 129
CttttcttcccttagctgttccTTTCCTGTGGTTTTAAAAAAGTGACCAGAAACTAGGTCTCT
ATTTTCATTGCTTTGCTGCATATTCTTTTAACCTGCTTTTATCTTTTACAGAGTT
GAGGGGCTTT**Y**TAAATAACCTAGACAATGTCAAGATTCTTAGCTGCGTTTTCT
GTCTAAAAGTGTAGATGTCTAGTTATTCCTCATGTAAAACACAACATTTCAAC
CCTGAGTACTATAAACTTTATTATGCTTCTAGGTTACTTTTTCTCTTTAAGCAA
TTATTCCTACATTCCTAAGTGTTCACCAGTGGAACAGATAAGAGATAGAAGT
AGTTAGAAATTGAGATAATTGggttgacctgtcattgttgc
For: 5'-3' = cttttcttcccttagctgttcc
Rev 5'-3' = gcaacaatgacaggtcaacc

M78 = B9.60a (301 bp) **C to T** at position 197
CttcaggcattatttttttggtTCTCCACTACAGGAGAAATGTAAATGTGATGAGTCAGAAT
TTAGGATGGCTGTATGGGTTTCTTTGACTAATACAAGAAATCACTTTGTAATG
AATGAAATCAGTGGTTTCTGCATTACTCCGTATGTTCGACATGAACACAAATT
GATACACTTAACAAAGATACTTCTTTC**Y**GCCCTTCCAAATATTTCAAAATAAG
CTGGTCATAGTACTTGCTTTTCATAAAAAGATGGTAAGCTTCCAATATTTAGA
TTTaaggaaaggtgaaggaacactat
For: 5'-3' = cttcaggcattatttttttggt
Rev 5'-3' = atagtgttccttcacctttcctt

M79 = B9.42 **Homopolymer in tree** (425 bp = majority men). A's. 8 A's to 9 A's (426 bp derived). Extra "A" inserted after position 212.
AgccagttggatgacacgttAAAAGTCATGCAATGGGAAAAAAAATTAAATAAATTGAC
AGAGAATTAAAAATTGTGGAAGTATGTCTCCAGAAGATGTGCCTACAGGGAA
AACAGAAGGACTCCTTCAGGCTGACATGAAAGGATATTACTGAGTAGTTCAG
AGCTACATAAAGAAAGTAATACCCCTGAGAAAGGCAACTAT**AAAAAAAA**TA
TAAAAGTTAGTATTACATATACAGCACGAGAGACAAAAAAAATATAGTTAGT
TCAGAACTAGAATCAGAAAGCAAGACAAATGGTGTTAATTAGATTGCTTGAT
GAGCTCATTATCATCAATATATTTTTCTTGTGAGACGAGGAATACTAGGAAAA
AAAAGGTACAAGTTAGAATTCATAAAATGTATAaaatgtcaggaaacgaagagg
For: 5'-3' = agccagttggatgacacgtt
Rev 5'-3' = cctcttcgtttcctgacattt

M80 = G10.107. **Homopolymer in tree** (290 bp = most men). 9 T's to 10 T's (291 bp derived). Extra "T" inserted after position 55.
Actttctcttcttttagggtgacc AATTAATTCTGATTTGCCTTGA**TTTTTTTTT**GGCATTTTT
ATGGCACCATAAAAACCATAAATGATTTGTATTCATTTTGGCAACCCTAGTTC
CAGGTTGATTGTGATGGCTGGTTGTGATGGCTATTTTGAAAGTTGGCTTTCCT
CTGTCCCAGATATTTTCTCTAAAACCTTTATAATTTTGTCTTATGGCTAGCTAC
ATAGAATTTTAAAATATTACAAATGGCCAGACAGTCCTACTTCAccataagattttgtgt
gtgtgt
For: 5'-3' = actttctcttcttttagggtgacc
Rev 5'-3' = acacacacacaaaatcttatgg

M81 = B9.58a (422bp) **C to T** at position 147.
ActtaatttatagtttcaatccctcaGTAATTTTAACTTACTTCTATTTTAAGAACTATAACCA
AACTATCTGTAAGACTTTTAAGCACTATCATACTCAGCTACACATCTCTTAAC
AAAAGAGGTAAATTTTGTCCTTTTTTGAA**Y**GTCATAGAGTATACTCACACAA
ACCAAGAAGAAACAATCTACTACATACCTACGCTATATG**G**TATATAACTATT
GCTCCTAGGCTACAAATTAGTGCGACACTA**T**TGTACTGAATATTATAGGCCAT
GTAACACAATGGTTTAAGTATCTGTGCCTCTAAACACAGAAAAGATATAGTG
AAAGTACAGTATTGCTCCTTTATTAAACTCAAAATGTTATGCAGCATATGACC
GACTATAAAATAGCGCTTATccagatacagacatctccatgaa
For: 5'-3' = acttaatttatagtttcaatccctca
Rev 5'-3' = ttcatggagatgtctgtatctgg

M82 = B9.t18 (328 bp ancestral). **Two bp deletion (-AT)** at position 179. (326 bp derived). This STS also contains **M69** which is normally associated with STS B9.62 at site a. The M82 deletion mutation is always linked to the M69 mutant C allele.
CtgtactcctgggtagcctgtTCAAATCCAAAAGCTTCAGGAGGCTGTTTACACTCCTGAA
A**T**AAAATATATTTCAGCAAGACAAAGGGAATAAAGATCCAAAAAAACAGGA
GAGCTAAGGGGAGATAAATTTTTCATGTTACATTCAATATCTCATGCAATAAT
TCTGCATTTTCAT**AT**GTTTCCAGGTAGGTTTGTTTCTTCAGTAGGTATTAAAC
ATTATTTTATAATCTTTCCTTACATGCTTCATGCCATTTGAATTATAGTCCCTT
GCCTCTGGTTCAGTCAAGTCTCTATCATTCTAgagttagtgtgttcaatcgttctt
For: 5'-3' = ctgtactcctgggtagcctgt
Rev 5'-3' = aagaacgattgaacacactaactc

M83 = B9. Alu01 (503 bp) **C to T** at position 120
GggaaaggagttatccagaaaAGTGAAATTTATTCTAAAATTTTAAGTTTCCATGTTTTA
AAGAGAGGCAGCAATGAGAAAAAAGGTTAAGAACAAGTAGGAAATACTGAA
ATAATGGG**Y**CAGGCACGGTGGCTCATGCTTGTAATCCCAGCACTTTGGGAGG
CCAAGGCAGGCAGATCACAAGGTGAGGAGATTGAAACCATCCTGGCTAACAT
GGTGAAACCCCATCTCTACTAAAAATACAAAAAAATTAGCCAGGTGTGGTGG
CACACACCTGTAGACCCAGCTACTTGGGAGGCTGAGGCAGGATAATGGCCTG
AACCCGGGAGGTGGAGCTTGCAATGAGCTGAGATCGTGCCACTGCACTCCAG
CCAGGGTGACAGAGTGAGACCCCGTCTCAAAAAAAAAAAAAAAGAATATTTG
AAATAATGTGTCTCTAAAATATGACAGACATGAGAATGAAGACAAAACATAA
GAAACTAAgctaagtaagcatgggtcatt

For: 5'-3' = gggaaaggagttatccagaaa
Rev 5'-3' = aatgacccatgcttacttagc

M84 = B9.72 **Homopolymer in tree**(439 bp = most men). 9 T's to 8 T's (438 bp derived). One deleted "T" at position 400.
CcctctccaactgagttcaagATGGAAACAGTTAAGACAGGAAAAATTCTATTCCATTTA
AACTCATATCATTAGAATCATAACTGCTTTCAGACCACAATATAATCACAAAC
CTGGGAAAATGGAAACTCATTAAGTATCAAAATACAAATCATATGCCACATA
TATTATATACCATTTTCAGCACTTGTCTCTTCTTAGAGGACACTGTAAAATAT
ATTTTATCATTGTTTAAAATAATTTGTTATATTTTGAAATTAAGCTCTATTACA
TTTTCCGTTTATTTTAAAGCTTTATTCTTACAAATTTTCTATACAGAGGTAAGT
TTTCTTCTATTTACATATATAAACATACATGTATACACAGAGAGACACAGTAA
CATATTTTATGC**TTTTTTTTT**ATTCCCACGGCAATTTCtggaagcagaaacgtatattgc
For: 5'-3' = ccctctccaactgagttcaag
Rev 5'-3' = gcaatatacgtttctgcttcca

M85 = B9.67a (568 bp) **C to A** at position 437
AacagaattatcaggaaaaggtttCATAAAATAAAAATCTTTTAAACTTATGAAAGATGCT
CAATATAAAAAACTGTAAACCAGGGAAATGCAAATAAAAATTACAATGAAA
TACTACACACCTCCCAGAATGGCTAAAATGAAAACAAAACTGTCAATTCTAA
GTGTTAGTGAGGACATGTGGTAACCAGAACTGGCATCCAATACTAGCTGATA
AACTCGTCAATCATTTGTAAAAACAGTCTGACAATAATCCACTAGTGAAAAT
ATACATAGTCTCAGTCACAGCAATTCTATCCTGTCTATCTAGGTA**A**CAGAAAT
GTCTACATACGTTACCTAGAAACATATACTTTAATATCCACAGAATTACTTGA
AATAGCCAAAAATTGGTAACTACCAAAAGTTGAATGGTAAAACAGATAGAA
AAAAAGCTATG**M**CTAACAAAACTACACTTAATAGAACACAAGC**G**TGAGCAT
TAATA**G**AACCATATAAATGCATTTTTTGAACCACTAAAAGAAGAAGCCAATA
CAAAAGAGGTGATTAAttgaaagtacacgaacaagtaaaa
For: 5'-3' = aacagaattatcaggaaaaggttt
Rev 5'-3' = gcaatatacgtttctgcttcca

M86 = B9.t25a (324 bp) **T to G** at position 85
TcccattatttgctatatttgctACATACATCTAAGGTCATATCAAAGAAAGAAAACACCAG
TCCAAGTGGTTAACACACAAGC**K**TATATAACTTGCTTCTGTCATAGATCAAG
TACTTCTGAGTAAGCTATTTTTTTGCGGTTAAATGTAATAAAAGCTTGTGTAT
GCCTAAACTATATTTAATAACAGCAGAACGTAGAAATATTTGAATCTTATATT
TTTGTCCCTACAGCAGTCAGATGTTTAGAACCCCGTGGAATGTGGCGATCTGA
TAC**T**AATATTCTGATGCCAGCTTGTTCgggtcagaaaagttaaatgagaaa
For: 5'-3' = tcccattatttgctatatttgct
Rev 5'-3' = tttctcatttaacttttctgaccc

M87 = B9.t25b (324 bp) **T to C** at position 277
TcccattatttgctatatttgctACATACATCTAAGGTCATATCAAAGAAAGAAAACACCAG
TCCAAGTGGTTAACACACAAGC**T**TATATAACTTGCTTCTGTCATAGATCAAGT
ACTTCTGAGTAAGCTATTTTTTTGCGGTTAAATGTAATAAAAGCTTGTGTATG
CCTAAACTATATTTAATAACAGCAGAACGTAGAAATATTTGAATCTTATATTT

TTGTCCCTACAGCAGTCAGATGTTTAGAACCCCGTGGAATGTGGCGATCTGAT
ACYAATATTCTGATGCCAGCTTGTTCgggtcagaaaagttaaatgagaaa
For: 5'-3' = tcccattatttgctatatttgct
Rev 5'-3' = tttctcatttaacttttctgaccc

M88 = B9.80 (314 bp) **A to G** at position 166
AttctagggtcaggcaactaggGAATACTGCTGTAGCCTAGAGCCTGCCAAAATTATTCA
AACTAGCCAATCCCATACTTCTTATCCTGCTCTGTCTTGCCTTTCCCTTGGTAA
ACCCAATATAGGCTATGGCCTAGGTGCTTTTCTTATTCCTGCTTCTTCTGCRT
ATCCAAGATAGGTTTTCCTCTCTAGCACTGTGTAGCATATAGTGACTACCTCT
CTAAGGCCTGTGATAATAATAAACTTTGCTTTCCTGAGTCTCTGTGGTCACAC
CTACTGACCATCACATggaagaccatagaatagaacaaaca
For: 5'-3' = attctagggtcaggcaactagg
Rev 5'-3' = tgtttgttctattctatggtcttcc

M89= B9.94 (527 bp) **C to T** at position 347
Agaagcagattgatgtcccact**T**AAAGAAGCAGTCTAGCCACATTTTGGTAGAGCAGCTG
TGGTGTGCCAGGGAGTCCCTTTCATCCCCTGGTCAGTTTTGTTTGCGCTCTCCT
AAACCTGCAGGCTGGAACAGCTGAGCCATCCAAACAGCAAGGATGACAACC
TTCCCTTTCTCCTAAGAACTCTGCCCCATTCAAGCTTGGCCCAACACTGTTGC
CAGGGGCTGGCTGGAATTCCAAGCTGGTGAGTCTTATCCTATGAGGTGCCAT
GAAAGTGGGGCCCACAGAAGGATGCTGCTCAGCTTCCTGGATTCAGCTCTCT
TCCTAAGGTTATGTACAAAAATCTYATCTCTCACTTTGCCTGAGTTGCAGCTA
CCTTTGCTGGTGATCCTGGACCCAAAGTGTGCCAGCCTCTCCTGATACTCTGT
GTGTACCTGAGCAGCTATTCTGCCAAGACTTCACACAGCTCTGTGCATGAAAC
CCAAGGCCTTAGTGAAGTGGGATCAtgaggggatctcctaactgga
For: 5'-3' = agaagcagattgatgtcccact
Rev 5'-3' = tccagttaggagatcccctca

M90 = B9.96 (331 bp) **C to G** at position 170
TgatgtttcttcagtctttgaggTTGCTGTCTTTTGGATTTTTGAAAAAATCCTATTTAATAA
CTTAGTGGGTTGGTTTGTAGCAACAGTGAATTCAATCAACTGGCTTTATTTCT
AGAATATTTTAAAGATATTTTATCTCAGGATTTCTGGATGGTGTTCTGTAACT
STAGGGACTGGGAATGAGCTTTGGCTTTGTTCCTTTACACCCTGAGGTTAGAA
ATCTGCTGCACTGGAGGGACCAAGATGCTCTCAGAGAAATGGTCACAACACT
CTAATGATTGGTAGTAGCCAATGTGCTTCATATGCGggtggtagcaggattcatctt
For: 5'-3' = tgatgtttcttcagtctttgagg
Rev 5'-3' = aagatgaatcctgctaccacc

M91 = B9.87a **Homopolymer.**(495 bp, most men = 9 T's). Either one T deleted or inserted at position 368 (i.e. 8 T's or 10 T's)
GagcttggactttaggacggGGAAAAGAAGTGCTAAATGTTTTTGAATAAAACCTTTACT
GCACATGATAAACATCCCTTAAAAATTACCTAGGAGCACCCTAAATTTTAAA
ATGATCACAAAGACCTGGACAGATTACAGTAAACCTTCAACATCGCTAAACA
CACGTACCATAAATCAAA**AGAAA**CACACTGCTAATGATCCGTTTTTTGATGT
GGAAATA**T**CATGCTGTTTTTAAGGGAAATTATACTTTATTGCGATGTTTTATT

TCAAAACAAGATGTTACACTTTATTTCCTATAATTTTATTTACAATATTTTACA
CCCGTTAAGCAAAAATCCCCCTACATTGCTATTCTG**TTTTTTTTT**AATCAG
TTCACTACTGTAGTATCTTTTTGTTCTCCATATATTTTTGAAAAATACGCAAAA
GGTAAGTTTTAAAAATCAAATGGTAGATTTTATTTGGAAGGGCACTgccagaagtg
ccttaaagttt
For: 5'-3' = gagcttggactttaggacgg
Rev 5'-3' = aaactttaaggcacttctggc

M92 = B9.G2 (470 bp) **T to C** at position 340
TtgaatttcccagaattttgcAATCTGATCCAAATAGTTCAATTTCACTCTAGTTTGGGCCT
GGGAAAGAGAGGGCCTTATAAGATTGGCATACTCCTTAACCTGACTTCATCG
AGTATGCAGTAAATGAACAAGTATTATTCTATGCTATCTACACTTCTCCACCA
ACGTGCCGGAGCCCCAGCTTCACTGTCTTATCTCACCAGCGGGGTCCACAAA
AAGCTCAAATAAGCTGAGTCTTTAATCTATAAAGAGCTAAGAATGTGCCGTC
TTAGGATCAACATCATGTCTAAATTTAAGGAATTATTCTTGGACTTAAAGGTG
GCTTGACCAAAAATA**Y**GTAGGCTCCAACAGTATTTAGACTCAATATCATCAA
GACACTCATTTAGAATGTACTGATATATAATTCAAAGAATTAAAATATTTTTC
TAGTTCATGTAAAAGAGCTggacacaaaaccagtttctgaa
For: 5'-3' = ttgaatttcccagaattttgc
Rev 5'-3' = ttcagaaactggttttgtgtcc

M93 = B9.93 (504 bp) **C to T** at position 459
AacaaaacaaaacaaaaatactgaaTCTTTAGAATTATGTACGCTAAGTGAAACATGTTTAT
AAACATAAATACACAGTTTTTATAAAATATTTTAAAGTTTTACGGATAATAAA
ACCTAAAAACTGGCCAGTCGTGGTGGCTCATGCCTGTAATCCCAACACTTTGG
AAGGCTGAGTCAGGTAGATCACGAGGTCAAAGGATCGAGACTATCCTGGCCA
ACATGGTGAAACCCCATCTCTACGAAAAATACAAAAATGAGTGGGCATAGTC
ACGCGCCTGTAGCCCCAGCTACTCAGGAGGCTGAGGCAGGAGAATCACTTCA
ATCCAGGAGGTGGAGGCCGCCTGGCCAGAGTGATAAGCTGCCTCAAAAACA
AAACAAACAAACAAACAAACAAAAACAATTAACTTATTATGTAAAATTACCC
TGCTAAATCAGTTTCCACACCCTGAGTTAAA**Y**CCAAGTCACACCAAGCTTTtaa
cctaaactatcttcaagtgaacc
For: 5'-3' = aacaaaacaaaacaaaaatactgaa
Rev 5'-3' = ggttcacttgaagatagtttaggtta

M94 = B9.122 (405 bp) **C to A** at position 227
CacatggagaacagagaaatgcAGTGCAGGGCAAGGGCCCACCCAGAAGCAACACAGTC
AATGGAGCCTCCTTCACCCAGGAAACTGCAAACTGAATGCATGATCCTAGGA
TCCTCTCCCATGGATCTTTGCAACTTTCAGGTCAGGAGATCCAGTCAGGGACC
CATTCCACTAGGGCCTTCAGTTAGAAACACAGAGCTCATGGAGTCTTATCAG
AGTAGCTGTT**M**AGGCATGCATAGGGACCCAGGAGCTTTATACACCCTGACCG
TAAAGTCCCCAGCAAATATGACTGAAATTCAAGCAAGGTGGAACACTAACCT
TTGCACATACACTTGGGAAGGGAGTGGAAATCAAGATGCCAAGCAGCATTGG
TCTGTGAACCccactttcacaacatttcacaag
For: 5'-3' = cacatggagaacagagaaatgc
Rev 5'-3' = cttgtgaaatgttgtgaaagtgg

M95 = B9.123 (480 bp) **C to T** at position 172
GagtggaaatcaagatgccaagCAGCATTGGTCTGTGAACCCCACTTTCACAACATTTCA
CAAGCTAAAAGCCCACTGGCTTGGATTTCCAGTCAGCTGCCAGCAATAGTGT
TGCACCTTCTTGGGATCAAATGGAGTTCCTGAGGATAAGGAAAGACTACCAT
ATTAGTG**Y**TGGATGGCTTAGCCTTTCCAACCTGTAGGCTTAGGAGAGTCCAG
ACTTACTAGGGATGTAAGGGATCCTCTTACACAAAACAGGTGCACTACCAAA
ATGTGGCCAGAGTGCTTTAAACAGGACCTTGACCCATTTCTCATCTCTGGGAA
GGACCTCACAACTGGGGCCTTCAAACACACCCACCCTCATTGTCTGGCTGAC
AAAGTTTTTACTTATTGCTGAAAAATAGTGCCCTGAGGGAAAGGCAGGCTCC
CATCACTGATGCTTTAATGACTCATCTGTTCTAGtctccaggttacagaaagccc
For: 5'-3' = gagtggaaatcaagatgccaag
Rev 5'-3' = gggctttctgtaacctggaga

M96 = G3.05a (440 bp) **G to C** at position 70. Internal lower case denotes location of alternative reverse primer region to amplify site a only, as 212 bp STS.
GttgccctctcacagagcacTTTAAAGTGAGCTGTGATGTGTAACTTGGAAAACAGGTCT
CTCATAATA**S**GATAAAACACTCAGGTATAATATTAAAAACCTATGGCAAAAT
ATATGGTCCTTTACAAAGCAACAAAGTGGGTGGGTGAATCTCTTCATTCTTGG
CTGGCCATCAGTTCCTGTTACTGTACaggagtgggaaaacagtagccCTGGGAAATGGGT
TAAAACTGAGTAGGCATCTCCTGTGTCCAATAAGAACTCAATATTTTTGTCTG
CTATATCAAGGGTTACTTGAGGCTCCTCTGTGGAGATGGTAAGTTGTCCAGTG
GGAGATATAGAGAATGTTAGGCC**T**TATAGGTTCTCTACTTTTTTGGCCATTAT
GAGTCTGAATGTCTCAAACTCCCTTTTTATCCTGGTgcaatccttccagtgacctt
For: 5'-3' = gttgccctctcacagagcac
Rev 5'-3' = aaggtcactggaaggattgc

M97 = G3.05b (440 bp) **T to G** at position 355
gttgccctctcacagagcacTTTAAAGTGAGCTGTGATGTGTAACTTGGAAAACAGGTCT
CTCATAATA**G**GATAAAACACTCAGGTATAATATTAAAAACCTATGGCAAAAT
ATATGGTCCTTTACAAAGCAACAAAGTGGGTGGGTGAATCTCTTCATTCTTGG
CTGGCCATCAGTTCCTGTTACTGTACAGGAGTGGGAAAACAGTAGCCCTGGG
AAATGGGTTAAAACTGAGTAGGCATCTCCTGTGTCCAATAAGAACTCAATAT
TTTTGTCTGCTATATCAAGGGTTACTTGAGGCTCCTCTGTGGAGATGGTAAGT
TGTCCAGTGGGAGATATAGAGAATGTTAGGCC**K**TATAGGTTCTCTACTTTTTT
GGCCATTATGAGTCTGAATGTCTCAAACTCCCTTTTTATCCTGGTgcaatccttccagt
gacctt
For: 5'-3' = gttgccctctcacagagcac
Rev 5'-3' = aaggtcactggaaggattgc

M98 = G3.04a (395 bp) **G to C** at position 158; has (GTTTT)6 motif
GaatggggtgttacatggagaCTACAGGGCTGTTATATTCATAACTTTAGGCTATCATTAT
TGAGGGCTGGATGTCCCTCTGAGCCTCAGGATTC**AAA**GGATACT**GTTTTGTT**
TTGTTTTGTTTTGTTTTGTTTTTTCCCA**C**GGGTAATTAACACTG**S**GTTTTAGG
ACAGTCTGGACTGGGGGTACATTAACAGTTGTACTAGAAACTTCCATGTCTCA
AACAGAGGGGTCTACTAGAGAAGCAATATGTCATGGAAGGCAGTTCTTCTCC

ATATCTGTGTAAAGGCAAGTATTTGAAGCTAGGAGAACTGTTCCTTCTGGCCT
GTTGCCCTCTCACAGAGCACTTTAAAGTGAGCTGTGATGTGTAACTTggaaaacag
gtctctcataatagg
For: 5'-3' = gaatggggtgttacatggaga
Rev 5'-3' = cctattatgagagacctgttttcc

M99 = G3.04b (395 bp nominal) **1 bp deletion** (3A's to 2A's) at position interval 96-98 , STS alos has polymorphic (GTTTT) motif
GaatggggtgttacatggagaCTACAGGGCTGTTATATTCATAACTTTAGGCTATCATTAT
TGAGGGCTGGATGTCCCTCTGAGCCTCAGGATTC**AAA**GGATACT**GTTTTGTT**
TTGTTTTGTTTTGTTTTGTTTTTTCCC**A**CGGGTAATTAACACTG**G**GTTTTAG
GACAGTCTGGACTGGGGGTACATTAACAGTTGTACTAGAAACTTCCATGTCTC
AAACAGAGGGGTCTACTAGAGAAGCAATATGTCATGGAAGGCAGTTCTTCTC
CATATCTGTGTAAAGGCAAGTATTTGAAGCTAGGAGAACTGTTCCTTCTGGCC
TGTTGCCCTCTCACAGAGCACTTTAAAGTGAGCTGTGATGTGTAACTTggaaaaca
ggtctctcataatagg
For: 5'-3' = gaatggggtgttacatggaga
Rev 5'-3' = cctattatgagagacctgttttcc

M100 = G3.04c (395 bp nominal) **in tree (penta microsatellite)** (GTTTT)5; (GTTTT)6 = most men); (GTTTT)7; (GTTTT)8 alleles detected
GaatggggtgttacatggagaCTACAGGGCTGTTATATTCATAACTTTAGGCTATCATTAT
TGAGGGCTGGATGTCCCTCTGAGCCTCAGGATTC**AAA**GGATACT**GTTTTGTT**
TTGTTTTGTTTTGTTTTGTTTTTTCCC**A**CGGGTAATTAACACTG**G**GTTTTAG
GACAGTCTGGACTGGGGGTACATTAACAGTTGTACTAGAAACTTCCATGTCTC
AAACAGAGGGGTCTACTAGAGAAGCAATATGTCATGGAAGGCAGTTCTTCTC
CATATCTGTGTAAAGGCAAGTATTTGAAGCTAGGAGAACTGTTCCTTCTGGCC
TGTTGCCCTCTCACAGAGCACTTTAAAGTGAGCTGTGATGTGTAACTTggaaaaca
ggtctctcataatagg
For: 5'-3' = gaatggggtgttacatggaga
Rev 5'-3' = cctattatgagagacctgttttcc

M101 = A8.05a original (460 bp) **C to T** at position 154
Tcacagcagcttcagcaaa**C**ACAGATTTCTGGTGTTGGAGGACAGATTTAACTACAGAA
AATTCTGTTGGGCAATCGGAAGCCTCAATCTATACAGACTTTTAGGAGGAG**C**
CTGCCTGTTTGGTTCAAATTTAGCCAAAATA**TTTTTTTTTTA**Y**C**ACTGATTCA
GTAAATCTCCTAACTTTGCAGGAACTGGGATCCTAAAAATTATGGAACGAAT
TGTAGAAACTCAAGCAACTTTCTCCAAAGCCTAGGGttcagcaagagtaagcaagaggCA
CTGAGCCGCTGGAGTCTGCACATTGATAAATTTACTTACAGTCGTAAATAAAT
TGCATCATCTTCAgctagtaacacagagtctaatttttatAGCGGCATACTTGCCTCCACGACT
TTCCT**A**GACACCAGAAAGAAAGGC**G**AGAGCCAGCCTTAGCCTAATCaagaaccat
gatccaaaaagg
For: 5'-3' = tcacagcagcttcagcaaa
new R 5'-3' = ataaaaattagactctgtgttactagc3' (used with F primer, just amplifies (369 bp) the first 2 sites including homopolymer T region
Rev 5'-3' = ccttttggatcatggttctt

M102= B9.101 (480 bp) **G to C** at position 301
AaactgggacacttgtaatgaatAATTACTTTGTTTGTAAATCACAATAGAGATTCTCCATA
TCAAAGCTGTGAACTGTATTCTATAGTATTTAGGCAAATAAGATAGCTACAA
ATTTAAGTACTGTAATAATAGATGCCTGACAATATGTGCTATAGGTAAATCTT
TGAAATTTATTAAATGAAGTATAGATTGAATACAAGTAATATGTAATAATAC
ATTATAATTTAATAACATTTAGAATAATTACATTTTATACAAAAATAAAATTA
AGAtaaaattcacatagtgcaatggtgA**S**TAAGATGTGAAAAGACAATAAGAATAAACAGC
ATTAAAATTATTGATAGAGTTTGTAAAACCCCTAGAGATTAAGGAAAACAAA
CATAGGAATAAATTAGAAAACTAGAGACAATAATAATTTCTGTAAATTATAG
GCTACCAAAACCAGAATaagaataaacaaggactcaaaaaac
For: 5'-3' = aaactgggacacttgtaatgaat
New R 5'-3'-taaaattcacatagtgcaatggtg
Rev 5'-3' = gttttttgagtccttgtttattctt

M103 = B9.117new (463 bp) **C to T** at position 259
CagtaagtgaactcacacataattccACAGGCATCTGAGCCCGTAGCAGCCTCAGCTGCCAT
TTTGATGGCAACCTAGATACTGGGGTTCTACAGACACAACTGCAGCCACTGT
ACTGCTCCAAGGACACAGAACAGGTATACACACACACCCATGGAGGGGTATT
TGCCACATTGCTATGAGCTGCTGTTGAGACTGAGAATTGGCCAGACCATGCTC
TTCACAGCTTCTTGCTCCTGCTCCTTGCCTAGGTTCTCC**Y**CCACCTTCTCTGGT
CTTGAACCCAATATGCCATTTTAGAGAGTTTGATGTTGGATAGTACCCCACCC
TTGGCCTGAGTTCAGGTTGATGCAGTTGCAGTCGCTGCCCATCCAAGAAGAG
ACAAAAACACTAGGCTATCCTCTTCATACTTAGAATAATATCCACTGCTCTGC
AACAAGACgctgtgaaactgaaataaaactgg
For: 5'-3' = cagtaagtgaactcacacataattcc
Rev 5'-3' = ccagttttatttcagtttcacagc

M104 = DYS257a (288 bp) Duplicated locus. Most men have both **A and G** alleles at position 162, however some have only A allele. The second site at position 202 is often just C, although sometimes both **C and T** alleles occur.
GaacttgtcgggaggcaatGGTGACATTCATTGTGACCTTAGCCAGAGCTCACAATCAA
CCATGGTGCACTGAGACTAGCTCATGCACATTCATCAGGCAGATTCAGGCAC
CTGGCTGTCAGAGCTGTCAGCCTTCCTCAGTAGAGGAAAATGCTACAGTC**R**G
CACTGGCCTGGTATCAGGAAAATAGATGCCTGCAAAAA**Y**CCACTGTGGGACC
CTAAAAGTCTTGACCTCAGGTCCCCTTTGTGCTGTCTCTGTTGTCAGGATccacta
aaggaggaagtgtatca
For: 5'-3' = gaacttgtcgggaggcaat
Rev 5'-3' = tgatacacttcctcctttagtgg

M105 = B9.6-7a (572 bp) **C to T** at position 478
GggaggcaacctaagaaagGTGTACAACTGTCCTGACATTGGATTGCCTGCTTACTGTG
AAGTATGTGAACAATTTGTGACTCAGAACTTTAGTGAGATTTTTATAGGCAGA
AGTTCTCATCATGCCTCATCAGAATTTTCCGTTAACAAGTGTCAGAGAATCTG
TAATGGCTTGAGAATCATGACTTTCCTCCTATTTATGGAAGAGGAGAAAAAA
GAAATTTCGAAGACAATTCTCAGATTTAGATAAATTATCTCAGGATTTTCTAT
ATATTTTACCTGGTCCCTATGGTGTGGTAAGGTAAAGTACACTGTACTTGGAC

AGGTGAAGCAATTTCTACTCTACTAGGTCATCACCAAGCATAGCTTTGTTACT
GGGAAAGCTAATTATAGTTCCCTATGACAGTATC**A**AAGAAAGAAAGAGGTGA
AAAGAGTAGACAATAAGGAAGGTAGGTATGATTATAGGCATGAGAAATG**Y**T
ATGGGTAATAACGTGTTCTACACTGACTCAAGTCAGCAAGGAGTAGGTGGAA
AAGCGAGAGATTCAATCCAGGatgacagaatgcgttcacct
For: 5'-3' = gggaggcaacctaagaaag
Rev 5'-3' = aggtgaacgcattctgtcat

M106 = B9.6-7b (572 bp) **A to G** at position 411
GggaggcaacctaagaaagGTGTACAACTGTCCTGACATTGGATTGCCTGCTTACTGTG
AAGTATGTGAACAATTTGTGACTCAGAACTTTAGTGAGATTTTTATAGGCAGA
AGTTCTCATCATGCCTCATCAGAATTTTCCGTTAACAAGTGTCAGAGAATCTG
TAATGGCTTGAGAATCATGACTTTCCTCCTATTTATGGAAGAGGAGAAAAAA
GAAATTTCGAAGACAATTCTCAGATTTAGATAAATTATCTCAGGATTTTCTAT
ATATTTTACCTGGTCCCTATGGTGTGGTAAGGTAAAGTACACTGTACTTGGAC
AGGTGAAGCAATTTCTACTCTACTAGGTCATCACCAAGCATAGCTTTGTTACT
GGGAAAGCTAATTATAGTTCCCTATGACAGTATC**R**AAGAAAGAAAGAGGTG
AAAAGAGTAGACAATAAGGAAGGTAGGTATGATTATAGGCATGAGAAATG**C**
TATGGGTAATAACGTGTTCTACACTGACTCAAGTCAGCAAGGAGTAGGTGGA
AAAGCGAGAGATTCAATCCAGGatgacagaatgcgttcacct
For: 5'-3' = gggaggcaacctaagaaag
Rev 5'-3' = aggtgaacgcattctgtcat

M107 = B9.112n (376 bp) **A to G** at position 298
CaaaagcactcgggttcctTGTTTCAATCCCACCTCACATACACATAAGCATCATTAACA
GTACAGCGTGGGGCTCTTTATCCCATCTTGTGCACCGCTTGCCTGAGAGAATT
TGCTACTGGTCCTGGGGAGCCCTGTCATATTCCCTTAGCAGGCCTGCAAAGAT
CTGTGTCCATTTCTTTTCCAAAAAGTCATTTTTCTCTCAACATCCCAATCTCAT
TTCCAAAACTGTCAATAAATATCAAGTTTCTTAGATTTTACTCATTTCTTAAGC
CAACGTATTAACCTTCTAATTTC**R**TGAATGCTAATAGAAAGCATGAGACACC
TATGCATCATATAAAAGTGTTTTTTATTcgttgcataagtgggagtaaag
For: 5'-3' = caaaagcactcgggttcct
Rev 5'-3' = ctttactcccacttatgcaacg

M108 = B9.113n (321 bp)**T to C** at position 40. Probably **recurrent**
AgatggagccagcagaaagGAGAGAAGTAGATGAACATC**Y**GAAACTATACCTGAATG
TCAGAGAAAAGTGGATTGACTTCAGAGGAACAGCTTGATGGTGTAACTTTGG
AGAAGAATCCGGCTGGAGACTTTAGTGATCTGGGTAGAAGATAAAATCATCC
ACAATATTTACTGGGGTTTTTTTTGCATTTCCTGAATTTGAATCTTGGCCAGAG
TAAAGGGAAATATTCATCCCTCCTCCTTTTTAGCACCCATTCCCACTTAAAGC
CACCTCTATCACATAAAATCCTCCACATTTaccatcattcaattcatctgtgt
For: 5'-3' = agatggagccagcagaaag
Rev 5'-3' = acacagatgaattgaatgatggt

M109 = G3.15 (312 bp) **C to T** at position 264

GggtatcaaatgtcttcaacctAAAGTACAAGGAATTATTTCTCAGTGTTTGGAATGACTT
GACTTCCTTGAAAATATTGTTGCAGAGTTGGGGACTACTTTTAAAATATCCTC
CATTGAATGTAATTCTACATGAAAGCTTGATTTTTCAAGTGCAAAATGCAAGT
GAGAAATAAGGCATATCATTCATTAAACCCTAATTCCAGCACTTTTAAATGA
GCTACTTTCTTGTATAATATTTTAGCTATTAAGGAACAAATTGT**Y**GCTTAAGA
AATGTATCTATCTTAAAAATgcaagtagcaggaaattccc
For: 5'-3' = gggtatcaaatgtcttcaacct
Rev 5'-3' = gggaatttcctgctacttgc

M110 = B9.86n (389 bp) **T to C** at position 241
CagggaaggaccgtaaaaggCTGTGGTGCTGATCAACGAAGGATTTCTCGGAGAAAATT
CCTCCTTTGCGGAAATGTCCGTAGAAACGCACCTTTTTTTTTTCCTGCCAGGA
CAAACCGCCGGCGATATCCGTTCATGTGAAAGTGTTTACTAACATTCTCTGAA
GACTCACTGGGTTCTCAGCTCGAGAACGTTCCTGTCACAAGACGTTTAGGAG
GCAGGATGCCGGTACAATGTATT**Y**ATGTTCTTGTAAACTGTTGCATTAACAGT
GCACTTCAAGTGGGCACATTTGTCGTTGGATTTTTTACCAACTCGAGCTTGGA
CTTTAGGACGGGGAAAAGAAGTGCTAAATGTTTTTGAATAAaacctttactgcacatgat
aaacat
For: 5'-3' = cagggaaggaccgtaaaagg
Rev 5'-3' = atgtttatcatgtgcagtaaaggtt

M111 = G3.19 (393 bp) **-2bp (TT) deletion** at position 188-189 interval. Polymorphic STS = 391 bp.
AatcttctgcaaagggttccTTTGGGTTTTGTTGTTGTTGTTTCCAATGCTAGCCAGA
GCAATAATTCTGAAAGGAAACCAAATTCCAAAATACAATGCAGATCTTCGTA
ATATTGTATTGTAACACAGTGTATCTAACATAAACAGTATGCCAAAAACAAC
AGAACAAGTTCTGTTTTTCACA**TT**GTTTTCTCCCCAAAATTTACCTTTCACAC
AAAACAAGTACCACAAAGAAGTGTCACAGCCTAAGAAACTGCCTTAGTATAA
CATTAAGAGCTTACATCCAGATTTACATCTGATAAAATATGACTGCTGGTATT
AACTTTAGGGCATATAAGGTATCTTCATCTCTTCTGAAAGAAGTGGgtccagtatttt
gttttgtagctg
For: 5'-3' = aatcttctgcaaagggttcc
Rev 5'-3' = cagctacaaaacaaaatactggac

M112 = G3.17a (445 bp) **G to A** at position 286
ActttttccaacagttatttttgaACTTCACTGTTACACAGTTGAGGTGACATTCATTATAAA
GAATACACAGAGGCTACTATATTAACCATTATATCTATATCTTTAGTT**A**ACCT
GAACGAAGTTGAGTAGATAAAATAAGATTCACATTAGGTAAAAAAACAAAA
ACAAAAACAAAAACAAAAACAAAAAACACAAACTCTACAGAAGTCTTGAAA
AGCAAAAGAGAACTGCCTCTTATAAAATCATATCCTTAAAAAAGAGGTGAGA
TAAAAACAAAGCAGT**R**TTTTTATCAGTACTGCATCCTTTTTTTCA**CA**GTTATT
TTCATTTACAGTTTGAAAGAGGTAGATAATTCTGCAACAGACAAGAATTGAA
CTGTGATTATCAGGTGTAATAAAATAGTTCCATTAACTTAGAAATattggtctcatcat
caagaaatata
For: 5'-3' = actttttccaacagttatttttga
Rev 5'-3' = tatatttcttgatgatgagaccaat

M113 = G3. 17b (445 bp) **A to G** at position 112
Acttttttccaacagttatttttga**A**CTTCACTGTTACACAGTTGAGGTGACATTCATTATAAA
GAATACACAGAGGCTACTATATTAACCATTATATCTATATCTTTAGTT**R**ACCT
GAACGAAGTTGAGTAGATAAAATAAGATTCACATTAGGTAAAAAAACAAAA
ACAAAAACAAAAACAAAAACAAAAAACACAAACTCTACAGAAGTCTTGAAA
AGCAAAAGAGAACTGCCTCTTATAAAATCATATCCTTAAAAAAGAGGTGAGA
TAAAAACAAAGCAGT**G**TTTTTATCAGTACTGCATCCTTTTTTTCA**C**AGTTATT
TTCATTTACAGTTTGAAAGAGGTAGATAATTCTGCAACAGACAAGAATTGAA
CTGTGATTATCAGGTGTAATAAAATAGTTCCATTAACTTAGAAATattggtctcatcat
caagaaatata
For: 5'-3' = acttttttccaacagttatttttga
Rev 5'-3' = tatatttcttgatgatgagaccaat

M114 = G3.23 (434 bp) **T to C** at position 387
Ttaccacacagttgagtagttctaaa**A**AAACAGAGATATGGTAGAAAAAGGAGAGGAAATT
TTCATTACAAAATCAATAGTTACAACTAAAAGAGAAACATGTACACAAAATA
TATCCATCAGTACAATGATCACACTTAATCTTAATCAATGCCTAGAGGAGATC
CTGTGGAGAGGGCTTTTGAGTAGCATTTTACTTCATTCATTCCTTTGGGGTCA
GCCTCCAGATGGACTCCTGGGGCTCTTTTAGAGGAAGTGTTCAGCATATTGGA
AGAATCCAGGTCAGCACAGGAATGCGTCACAGGCACTGCTAAATCTACATCT
GCTACTTTCACAGAGACCTGCCCTTTCAGAATTCCCAGTTTCTCACTGAGTTC
ATTCCTTTC**Y**ATTTGAAGAGCCTTGTACAGCTTCTCtaaccgctccaattttatttg
For: 5'-3' = ttaccacacagttgagtagttctaaa
Rev 5'-3' = caaataaaattggagcggtta

M115 = G3.22 (413 bp) **C to T** at position 201
agtttacagtcacatcaatttgga**A**AGTCATACAAATATTGTCAAAAAACTGATCTGAATCA
AATATGCCATGCTTGTTTCTTAATCCATTGAAGTTTTACTTATCATTTAAATGA
CTTGACAATATTAGTCAGTTTATATTTTCTTTTATGTAGATATTATGGGCTCCA
GAGTTTAAATTAGTATTTGATTTCACATTA**Y**GAAACCATTATAAAAAAGTCTC
AAATTAAGATAATTTAAGGTGATGAACACACAAACGTACACTTTGAAAGGAG
AAGGCAATGAAAACATGCATTCCAATAAAGGGGGAAAATGAGGCTGATGTG
CAACATAGTTGGGGAAATTGGTAAGAAGCTTTCTGTTACCACACAGTTGAGT
AGTTCTAAAAaaacagagatatggtagaaaaagga
For: 5'-3' = agtttacagtcacatcaatttgga
Rev 5'-3' = tccttttctaccatatctctgttt

M116 = G3.25a (429 bp) Three alleles. **A to T** (M116.2) or **A to C** (M116.1) at position 176
aagtatgacttatgaagtacgaagaaa**A**TCAAGGCTATTAATCAAAAATACCAGCAAAACTTT
TCCTATAGAAGCAAAGATAATGTTATAATTGTTAATTTCTTTTTTATATAAAA
TAACTCACCAAAGGAATGCACATCT**ATCT**GCTTTCTGAAAAAATAATTTCAA
ACTGATA**H**CTGTCAATTTTAATTATCTTAATTAAAATAAGCCATATTATGTTT
TTCTATCATCTAATAAGCTCTTTAGTGAAGAGCTAAAAATATATATAAAGAAC
ATAAAATCATATCCAACTATTAAGGGAAGATGCTATTTTCATCTACTTGCAGT

TTTTCTACCCAAATATAAATAATTTGTTTTAGCCATATTATCTCATTACTGAAG
TATCATAGGATGACTGAGTAGACtgctcattgtaaaatctaactgaat
For: 5'-3' = aagtatgacttatgaagtacgaagaaa
Rev 5'-3' = attcagttagattttacaatgagca

M117 = G3.25b (429 bp) **-4bp deletion** at position interval 142 to 145
AagtatgacttatgaagtacgaagaaaATCAAGGCTATTAATCAAAAATACCAGCAAAACTT
TTCCTATAGAAGCAAAGATAATGTTATAATTGTTAATTTCTTTTTTATATAAA
ATAACTCACCAAAGGAATGCACATCT**ATCT**GCTTTCTGAAAAAATAATTTCA
AACTGATA**A**CTGTCAATTTTAATTATCTTAATTAAAATAAGCCATATTATGTT
TTTCTATCATCTAATAAGCTCTTTAGTGAAGAGCTAAAAATATATATAAAGAA
CATAAAATCATATCCAACTATTAAGGGAAGATGCTATTTTCATCTACTTGCAG
TTTTTCTACCCAAATATAAATAATTTGTTTTAGCCATATTATCTCATTACTGAA
GTATCATAGGATGACTGAGTAGACtgctcattgtaaaatctaactgaat
For: 5'-3' = aagtatgacttatgaagtacgaagaaa
Rev 5'-3' = attcagttagattttacaatgagca

M118 = G3.29 (478 bp) **A to T** at position 109
AttctaagtttcacttcctgatccACCACAGAAATCACTTTACAATGTTCTTCCCTTCCTCCA
TCACTGCATTCTTCTCAACCAGCTGACACTTGTGTTTTCTTTATA**W**GAGTAAG
TGGTATCTTTCTTTTGTTAGTAAAGTTTATCTCAGAAGCTCCTATGGTAAAAG
CAGCAGTAACCAAAGCAGAAGTTTCACATTAAAAGAAAACAAAGTTGTTGTC
CTTAATTTCAAGGGAATCAGCACATGGTAGCTGAATTCTCTCAATTAAGACTG
ATGTGTAGCTCAGCTCAGGTGTGGACAGTAGAGCTGAGACCTCCTGCTCCTG
AAGTATATGAAAAAATGTCCCCGAGTTTTCTGGAGAAATGATAAATTACACT
AATCCATCAGATTATTTTATATACTGTCAGTCCCAAAGTAGCTCAAGAATCTG
AAAGGAAATCAGTGTAAGAGCTAgaggtagcgtaatttagggaacta
For: 5'-3' = attctaagtttcacttcctgatcc
Rev 5'-3' = tagttccctaaattacgctacctc

M119 = G3.32 (330 bp) **A to C** at position 224
GaatgcttatgaatttcccagaCACAGCTACTGTACTATCTCCAATCAGCACATTTTAAAG
AAATCTTAACTTAAATAGGGAAATGCCAAGGTAAATGACTCACCCTAAGGAA
GTCACGAAGTGCAAGTTAGAGATCTCAGTTTCAGAGTTTATGCTCCAAACCG
CAGTGCTATGTGTTTATTTGGGGAGACAGATAATTCTGCTCTTTAAAATTGCT
ATTTT**M**GCCTGTATGCTGAATTGGAATAACCCATAACATTTTTCTACATCTA
ATTTTAAAAAACGGTTTAAATTTTGTATTAATTaagaatacatcttgtatattgtgtgaa
For: 5'-3' = gaatgcttatgaatttcccaga
Rev 5'-3': ttcacacaatatacaagatgtattctt

M120 = B9.87b (495 bp) **T to C** at position 224
GagcttggactttaggacggGGAAAAGAAGTGCTAAATGTTTTTGAATAAAACCTTTACT
GCACATGATAAACATCCCTTAAAAATTACCTAGGAGCACCCTAAATTTTAAA
ATGATCACAAAGACCTGGACAGATTACAGTAAACCTTCAACATCGCTAAACA
CACGTACCATAAATCAAA**AGAAA**CACACTGCTAATGATCCGTTTTTTGATGT

GGAAATA**Y**CATGCTGTTTTTAAGGGAAATTATACTTTATTGCGATGTTTTATT
TCAAAACAAGATGTTACACTTTATTTCCTATAATTTTATTTACAATATTTTACA
CCCGTTAAGCAAAAATCCCCCTACATTGCTATTCTG**TTTTTTTTT**AATCAGTT
CACTACTGTAGTATCTTTTTGTTCTCCATATATTTTTGAAAAATACGCAAAAG
GTAAGTTTTAAAAATCAAATGGTAGATTTTATTTGGAAGGGCACTgccagaagtgcc
ttaaagttt
For: 5'-3' = gagcttggactttaggacgg
Rev 5'-3': aaactttaaggcacttctggc

M121 = B9.87c (495 bp) **5 bp deletion** at position interval 183-187
Gagcttggactttaggacgg**GGAAA**AGAAGTGCTAAATGTTTTTGAATAAAACCTTTACT
GCACATGATAAACATCCCTTAAAAATTACCTAGGAGCACCCTAAATTTTAAA
ATGATCACAAAGACCTGGACAGATTACAGTAAACCTTCAACATCGCTAAACA
CACGTACCATAAATCAAA**AGAAA**CACACTGCTAATGATCCGTTTTTTGATGT
GGAAATA**T**CATGCTGTTTTTAAGGGAAATTATACTTTATTGCGATGTTTTATT
TCAAAACAAGATGTTACACTTTATTTCCTATAATTTTATTTACAATATTTTACA
CCCGTTAAGCAAAAATCCCCCTACATTGCTATTCTG**TTTTTTTTT**AATCAGTT
CACTACTGTAGTATCTTTTTGTTCTCCATATATTTTTGAAAAATACGCAAAAG
GTAAGTTTTAAAAATCAAATGGTAGATTTTATTTGGAAGGGCACTgccagaagtgcc
ttaaagttt
For: 5'-3' = gagcttggactttaggacgg
Rev 5'-3' = aaactttaaggcacttctggc

M122= G3.27a (393 bp) **T to C** substitution at position 73
TggtaaactctacttagttgcctttTGGAAATGAATAAATCAAGGTAGAAAA**G**CAATTGAGA
TACTAATTCA**Y**GCTCTCAGGGGAAAATCTGAATAAAGCTATCTTTTCTAACA
CAGAGCAAGTGACTCTCAAAGTCACAGTATCTGAACTAGCATATCA**G**CATCG
CCTGAATACCTAGAAATGCAAATTCCTGGGCAACACCAGAATCTAACAAAGC
AAAAAACTATGGGGGGAACAGGGAAGT**CG**GTTTAATAATACTGAGTTTGTGC
AACCTCAACTTTGCTTTA**T**AGGAAAGCAAAATCTCAATATGATAA**A**GTTTTCT
TCAACAAAACTCTGAGATAACTATGTTGAGGGAAAGAAGTTGATCACATgcaag
aaaatctaattcgctg
For: 5'-3' = tggtaaactctacttagttgcctttt
Rev 5'-3' = cagcgaattagattttcttgc

M123 = G3.27b (393 bp) **G to A** at position 161
TggtaaactctacttagttgcctttTGGAAATGAATAAATCAAGGTAGAAAA**G**CAATTGAGA
TACTAATTCA**T**GCTCTCAGGGGAAAATCTGAATAAAGCTATCTTTTCTAACAC
AGAGCAAGTGACTCTCAAAGTCACAGTATCTGAACTAGCATATCA**R**CATCGC
CTGAATACCTAGAAATGCAAATTCCTGGGCAACACCAGAATCTAACAAAGCA
AAAAACTATGGGGGGAACAGGGAAGT**CG**GTTTAATAATACTGAGTTTGTGCA
ACCTCAACTTTGCTTTA**T**AGGAAAGCAAAATCTCAATATGATAA**A**GTTTTCTT
CAACAAAACTCTGAGATAACTATGTTGAGGGAAAGAAGTTGATCACATgcaaga
aaatctaattcgctg
For: 5'-3' = tggtaaactctacttagttgcctttt
Rev 5'-3' = cagcgaattagattttcttgc

M124 = G3.27c (393 bp) **C to T** at position 246
Tggtaaactctacttagttgcctt**T**GGAAATGAATAAATCAAGGTAGAAAA**G**CAATTGAGA
TACTAATTCA**T**GCTCTCAGGGGAAAATCTGAATAAAGCTATCTTTTCTAACAC
AGAGCAAGTGACTCTCAAAGTCACAGTATCTGAACTAGCATATCA**G**CATCGC
CTGAATACCTAGAAATGCAAATTCCTGGGCAACACCAGAATCTAACAAAGCA
AAAAACTATGGGGGGAACAGGGAAGT**Y**G**G**TTTAATAATACTGAGTTTGTGC
AACCTCAACTTTGCTTTA**T**AGGAAAGCAAAATCTCAATATGATAA**A**GTTTTCT
TCAACAAAACTCTGAGATAACTATGTTGAGGGAAAGAAGTTGATCACATgcaag
aaaatctaattcgctg
For: 5'-3' = tggtaaactctacttagttgcctttt
Rev 5'-3' = cagcgaattagattttcttgc

M125 = B9.108a (367 bp) **T to C** at position 301
Gccaccctcttatgcctct**G**GCCTTTACAAAGACAGCTGGTAAGAGGCTGCCCAGCTCAT
CTGAAGTACAGGATAAGATTGTCTGACTTGGAGATACCATTTTCCACTTAGCA
GCCATGTAATCTTTCATATTCATTTTTTCTAAGTGGCACTTTTCTCAGATGTAA
AATGGGGATAATGAGTTTATTCATCTTTGAGTTGCTCCCAAGCAGAAGTCAAC
TTGAGACTATAAACTTGTGCTCACTGCAGTGCTTGAAACCGAGTTTGTACTTA
ATA**AATA**GCTGCATACATCTTTTTCTA**Y**ACATGTCAGATGCTTAATTGTGTTT
CCCGAAGATGTTGCCAAGCCgggtcctcacataactcctga
For: 5'-3' = gccaccctcttatgcctct
Rev 5'-3' = tcaggagttatgtgaggaccc

M126 = B9.108b (367 bp nominal) **4 bp deletion** (AATA) at interval 277-280.
Gccaccctcttatgcctct**G**GCCTTTACAAAGACAGCTGGTAAGAGGCTGCCCAGCTCAT
CTGAAGTACAGGATAAGATTGTCTGACTTGGAGATACCATTTTCCACTTAGCA
GCCATGTAATCTTTCATATTCATTTTTTCTAAGTGGCACTTTTCTCAGATGTAA
AATGGGGATAATGAGTTTATTCATCTTTGAGTTGCTCCCAAGCAGAAGTCAAC
TTGAGACTATAAACTTGTGCTCACTGCAGTGCTTGAAACCGAGTTTGTACTTA
ATA**AATA**GCTGCATACATCTTTTTCTA**T**ACATGTCAGATGCTTAATTGTGTTT
CCCGAAGATGTTGCCAAGCCgggtcctcacataactcctga
For: 5'-3' = gccaccctcttatgcctct
Rev 5'-3' = tcaggagttatgtgaggaccc

M127 = G3.30 (412 bp) **C to A** at position 372 bp
TgaaaggaaatcagtgtaagagcTAGAGGTAGCGTAATTTAGGGAACTAATCAGGAAAGA
GGTATTAACATTTCTGAATCCTTAGTTTCACTTATCCTTTCAATTCACAAGATT
GCTTTATTTCACATTTTGATAAAGACCAAAATGGTCCAAAAATAAGGGGAGG
AAGAACCTATACTACAAGAACCGAATTCCCAGACACTCAGGATAAACTTTAG
GTATATCCTTCAATCAGCTTTGTTCCAAATACAGGTAACGAGCCAGGCAATGT
TACGGAAAATAAGGGTAAGATAAAGCAAATATCCTGTGCTTTGGTTAACAAA
CAAAACTGTATCACAAGTCAAACTCGTACAAAAGGCAGGAGAAGAGGT**M**TG
GAAGATCTGTTAGGtgctgaactacagtcacctttaca
For: 5'-3' = tgaaaggaaatcagtgtaagagc
Rev 5'-3' = tgtaaaggtgactgtagttcagca

M128 = G3. 17c (445 bp vs 443 bp) **-2 bp deletion** (CA) at position interval 316-317
Actttttccaacagttatttttga**A**CTTCACTGTTACACAGTTGAGGTGACATTCATTATAAA
GAATACACAGAGGCTACTATATTAACCATTATATCTATATCTTTAGTT**A**ACCT
GAACGAAGTTGAGTAGATAAAATAAGATTCACATTAGGTAAAAAAACAAAA
ACAAAAACAAAAACAAAAACAAAAAACACAAACTCTACAGAAGTCTTGAAA
AGCAAAAGAGAACTGCCTCTTATAAAATCATATCCTTAAAAAAGAGGTGAGA
TAAAAACAAAGCAGT**G**TTTTTATCAGTACTGCATCCTTTTTTTCA**CA**GTTATT
TTCATTTACAGTTTGAAAGAGGTAGATAATTCTGCAACAGACAAGAATTGAA
CTGTGATTATCAGGTGTAATAAAATAGTTCCATTAACTTAGAAATattggtctcatcat
caagaaatata
For: 5'-3' = actttttccaacagttatttttga
Rev 5'-3' = tatatttcttgatgatgagaccaat

M129 = A8.04 (255 bp) **G to A** at position 221.
There is a polymorphic (CA)n motif immediately adjacent to the 3' end of STS
Aatggcttactacaaagaacatttc**T**GTAGTATATTTTTATGTATGTATGTATTATGTATTTAT
TTATTTATTTATTTTTGAGACAGAGTCACAATGCTGCCCAGGCCCTAGTGCAG
TGGTGTGATCTTAGCTTACTGCAACATCTGCTTCTGTGTTCAAGAGATTCTCCT
GCCTTAGCCTGTGGAGTAGCTGGAATTACAGGTGCACACCACCAAGCCC**R**GC
TAATTTTTAtcttctttggtagagaccgtgta
For: 5'-3' = aatggcttactacaaagaacatttc
Rev 5'-3' = tacacggtctctaccaaagaaga

M131 = A8.14n (306 bp) 9 bp **deletion** at interval 93 to 101
Cacacccagaatacaataatttt**A**AAAACATAATAAAGGTCAATTTAGAGCAGAGAAATTA
TTCTTTTAAATTACAAATGTTTGCTGTTCAG**GCAAATTAC**ACAGAAAGTTA
AGAATAACCCTTTAAATGATAGGAAAAGGCATTAGTAAGATAAAATGTGATT
ACTATTGAGATAAATATTTGCTATAAAAATAATTCAATTTGGTTAAACACAAA
TTGACTTCTTAAATAATCTTAAACATTAAGTAGAAGTAATTTTAGCTTATCAG
TAAATTTGAgaaaatgtacacttgtagaataaaaag
For: 5'-3' = cacacccagaatacaataatttt
Rev 5'-3' = ctttttattctacaagtgtacattttc

M132 = B9.67b (568 bp) **G to T** at position 482
Aacagaattatcaggaaaaggttt**C**ATAAAATAAAAATCTTTTAAACTTATGAAAGATGCT
CAATATAAAAAACTGTAAACCAGGGAAATGCAAATAAAAATTACAATGAAA
TACTACACACCTCCCAGAATGGCTAAAATGAAAACAAAACTGTCAATTCTAA
GTGTTAGTGAGGACATGTGGTAACCAGAACTGGCATCCAATACTAGCTGATA
AACTCGTCAATCATTTGTAAAAACAGTCTGACAATAATCCACTAGTGAAAAT
ATACATAGTCTCAGTCACAGCAATTCTATCCTGTCTATCTAGGTA**A**CAGAAAT
GTCTACATACGTTACCTAGAAACATATACTTTAATATCCACAGAATTACTTGA
AATAGCCAAAAATTGGTAACTACCAAAAGTTGAATGGTAAAACAGATAGAA
AAAAAGCTATG**C**CTAACAAAACTACACTTAATAGAACACAAGC**G**TGAGCATT
AATA**K**AACCATATAAATGCATTTTTTGAACCACTAAAAGAAGAAGCCAATAC
AAAAGAGGTGATTAAttgaaagtacacgaacaagtaaaa

For: 5'-3' = aacagaattatcaggaaaaggttt
Rev 5'-3' = ttttacttgttcgtgtactttcaa

M133 = A8.08F-newR (211 bp nominal vs 210) **1bp (T) deletion** at position 116. Site a. STS contains homopolymer A which normally has 10 A's, but sometimes 11 A's (sited).
TgaaatggaaatcaataaactcagtTTCCTCAAAGTTCAAAATACATGAGACTGCCTACCCT
CCTTGGAAGGCAAG**G**TGGGGCTTTCTGAAGCAAATACCAGCTTTAAAAAAAAA
A**T**GTATATATATATGAAGATATATAC**AAAAAAAAAA**TTTCCCCACAACCAGA
CAATCAGAATCATCAAACCCAgaagggttaaagaaaaagaaaagg
For: 5'-3' = tgaaatggaaatcaataaactcagt
Rev 5'-3' = cctttctttttctttaacccttc

M134 = A8.08newF-R (232 bp nominal vs 231) 1bp **deletion** (G) at position 54 (site b).
Agaatcatcaaacccagaagg**G**TTAAAGAAAAAGAAAAGGCCCAGGAAAGTAT**G**ATTG
GTGGGGATCAAAAGTATCTCTCCACAGTGGTAAATGAGAATTCTCAAAAAGA
GTAAAATTATAATTCTCATGCACATATAAAATAAATATGTATTACAGATTTTA
CTTAAACCATATAGCTCAAAATTAGCTAACAAGGAAGACATTATAACctgttcaaa
gagaagccaaaga
For: 5'-3' = agaatcatcaaacccagaagg
Rev 5'-3' =tctttggcttctctttgaacag

M135 = A8.08F-newR (211 bp nominal vs 212) **1 bp insertion** (+ C) at position 150 = site c, within homopolymer A track.
tgaaatggaaatcaataaactcagtTTCCTCAAAGTTCAAAATACATGAGACTGCCTACCCTC
CTTGGAAGGCAAG**G**TGGGGCTTTCTGAAGCAAATACCAGCTTTAAAAAAAAA
TGTATATATATATGAAGATATATAC**AAAAAAAAAC**ATTTCCCCACAACCAGA
CAATCAGAATCATCAAACCCAgaagggttaaagaaaaagaaaagg
Site a $(A)_{10}$ -TTT most males
Site c $(A)_9$CATTT = M135
Site d $(A)_{11}$TTT
For: 5'-3' = tgaaatggaaatcaataaactcagt
Rev 5'-3' = cctttctttttctttaacccttc

M136 = B9.61 (339 bp) **C to T** at position 196
Atgtgaagacaacactgtgtgg**G**AGAACCTAGGAAAGTAATTTTACATGCTAAAATGAGT
TTCCCTAGTTAATGTTAACATGAACTACCAACCGTATTACCTTCTCCTCAGGA
GATAAGTTTTGTTTGCTATTGCTGACAGGAAAGCCACTGCCAAATTCTTTGGA
ATGAATATCAGCTCCATATTCAACTGTCA**Y**GTCTTCCTCAATGCTGCTCACCA
GCCTCCAGAATTCCTTCTCTACAAGTTCTGTAGGCACCATCTGTGAAAACACA
TGTAAAAGGTTATCATAGCCCACTATACTTTGGACTCATGTCTccatgagaactaagac
taccacaa
For: 5'-3' = atgtgaagacaacactgtgtgg
Rev 5'-3' = ttgtggtagtcttagttctcatgg

M137 = G3.27d (393 bp) **T to C** at position 289

Tggtaaactctacttagttgcctttg**G**AAATGAATAAATCAAGGTAGAAAA**G**CAATTGAGA
TACTAATTCA**T**GCTCTCAGGGGAAAATCTGAATAAAGCTATCTTTTCTAACAC
AGAGCAAGTGACTCTCAAAGTCACAGTATCTGAACTAGCATATCA**G**CATCGC
CTGAATACCTAGAAATGCAAATTCCTGGGCAACACCAGAATCTAACAAAGCA
AAAAACTATGGGGGGAACAGGGAAGT**CG**GTTTAATAATACTGAGTTTGTGCA
ACCTCAACTTTGCTTTA**Y**AGGAAAGCAAAATCTCAATATGATAA**A**GTTTTCTT
CAACAAAACTCTGAGATAACTATGTTGAGGGAAAGAAGTTGATCACATgcaaga
aaatctaattcgctg
For: 5'-3' = tggtaaactctacttagttgcctttg
Rev 5'-3' = cagcgaattagattttcttgc

M138 = A8.17(442 bp) **C to T** at position 291
Aacttccaaaactgtgaaaagattg**T**TTTTAAAAGGCTATAACAGTGACTTTCAGGTGAAGA
CTTGGACAAATAGATAATTTCTGTACCCATTAAAATCAGGGGCTGTTACTATG
TTTGAAGACATTGTCGCCACAGCTTGAAGTCTGTAAGGAAAACCTGTAAAAT
TAGTGGGTGCCCACTCTAGTTTTAATCATTTGAGTTTCCACTCCTCATTGTGGT
TGAACTATTTTATAACTCTGCAAAATCTAGAAAGTTGAAAAGAAACCAAAGA
TACTTTCCCTTTTCTTC**Y**CACTTCTCCTACCCTTGGCCCACCTCCTTCTCCACC
TACTACTCCACATGGAACCTGGAGATTTGAGTCGGGGAGTGATGTAATACCT
GCGGCGCGTTGGCCCTTTACACACCTGTCAGCCATTTCAAGGCctgaaggggctgcttt
aatc
For: 5'-3' = aacttccaaaactgtgaaaagatt
Rev: 5'-3' = gattaaagcagccccttcag

M139 = A8.28a (459 bp nominal vs 460) **1 bp deletion** at position 401. **5 G's to 4 G's** .
Ttactgataatgccatattgttttg**G**CTTAATATCAGGCTAAGTAACCACAGTATTCTGATTT**A**
AAAAAAAACATACTAGAGAGCAAGTTTATTGACAAATCTTTAGGAACTTCAG
GTACAGCATATGATTTCTGAACTATGTGTGTAAATAAGGTTTTGTTTATTCAA
ATTTAACACAGGGTAGTCTGTGTATGCCTTCCGATTTGATAGCTCTAATAAAA
CACTTTAATAGTACCATATCAAATAAATTTTATCATCATCGATTTTCTTCTTAA
TATGAAATAACACATATTTGTGATTTTTCTAAGAGTCAAAATCTCAAAAATCA
TTTTAGGTATAAAATATACCCCGAAAGTTTTATTTTATTCCATTTTATAATTAA
TCTGACTTGGAAA**GGGGG**AAAAAAGCTCAAAGGGTATGTGAACATTTCATT
AAGATaggaccattggtgtctgagaa
For: 5'-3' = ttactgataatgccatattgttttg
Rev 5'-3' = ttctcagacaccaatggtcct

M140 = A8.28b (459 bp nominal vs 460) **1 bp insertion** within 9 A's **homopolymer** (most men) to 11 A's at position 73. **Recurrent** because 11 A's found in different haplogroups.
Ttactgataatgccatattgttttg**G**CTTAATATCAGGCTAAGTAACCACAGTATTCTGATTT**A**
AAAAAAAAACATACTAGAGAGCAAGTTTATTGACAAATCTTTAGGAACTTCA
GGTACAGCATATGATTTCTGAACTATGTGTGTAAATAAGGTTTTGTTTATTCA
AATTTAACACAGGGTAGTCTGTGTATGCCTTCCGATTTGATAGCTCTAATAAA
ACACTTTAATAGTACCATATCAAATAAATTTTATCATCATCGATTTTCTTCTTA

ATATGAAATAACACATATTTGTGATTTTTCTAAGAGTCAAAATCTCAAAAATC
ATTTTAGGTATAAAATATACCCCGAAAGTTTTATTTTATTCCATTTTATAATTA
ATCTGACTTGGAAA**GGGG**AAAAAAGCTCAAAGGGTATGTGAACA**T**TTCATT
AAGATaggaccattggtgtctgagaa
For: 5'-3' = ttactgataatgccatattgttttg
Rev 5'-3' = ttctcagacaccaatggtcct

M141 = A8.30a (424 bp nominal) **T to A** at position 51. Locus also has **two homopolymer** T tracks which are both polymorphic. See next below.
Catcttaaaatacatttcatagctt**T**CAAACTCAAATATGAAAACAATT**W**GTTTTTTAGA**TT**
TTTTTTTTCTTTTTACTTCAAGTTCTTTATATTCTAGACTAACACTTTAGGGCA
GATATTGGAGGGTGTGTCTCTCTTGGTGCAACTATTGCCTTTGCTTCAAATGG
TGGCATATGGAGGAGGACACAACCTGTAGGAAGTGTTCAAGGAGTCTGGTAG
TGACACCTGCTCAATATTGCTAGTGATAAAACTGTAGCCACTGTATAGCAATA
TCTGCCTGTAGAATGTCATTTCCTTTGAGGGGTACA**TTTTTTTT**AGAGTTTCC
TATAACCTCTAGAGCTGAACTTCATAAAAATAGGTAAAGGTTGGCCTTAAAA
AGCCTACATTACACACTTTCaggatgctagacctaatagtaagc
For: 5'-3' = catcttaaaatacatttcatagcttt
Rev 5'-3' = gcttactattaggtctagcatcct

M142 = A8.30b,c (424 bp nominal vs 423) T to A, **also has Homopolymers**10 T's to 9 T's at position interval 61 to 72 & 8 T's to 9 T's at position interval 311-319 **in tree**
Catcttaaaatacatttcatagctt**T**CAAACTCAAATATGAAAACAATT**T**GTTTTTTAGA**TTT**
TTTTTTTCTTTTTACTTCAAGTTCTTTATATTCTAGACTAACACTTTAGGGCAG
ATATTGGAGGGTGTGTCTCTCTTGGTGCAACTATTGCCTTTGCTTCAAATGGT
GGCATATGGAGGAGGACACAACCTGTAGGAAGTGTTCAAGGAGTCTGGTAGT
GACACCTGCTCAATATTGCTAGTGATAAAACTGTAGCCACTGTATAGCAATAT
CTGCCTGTAGAATGTCATTTCCTTTGAGGGGTACA**TTTTTTTT**AGAGTTTCCT
ATAACCTCTAGAGCTGAACTTCATAAAAATAGGTAAAGGTTGGCCTTAAAAA
GCCTACATTACACACTTTCaggatgctagacctaatagtaagc
For: 5'-3' = catcttaaaatacatttcatagcttt (
Rev 5'-3' = gcttactattaggtctagcatcct

M143 = B9.50b (385 bp) **G to T** at position 246
Atgctataataactaggtgttgaag**A**TAAAATCAGTTTAATTTAAATAAGAGGATAAAAGAA
GTATGAGCAGAAAAAGGTTTTCAATATTAACTAGGAAAGTCTGAAAAATAAT
CAGAAATTCTAAAGATAAAAACATAACATTAAAAATTATAAACTAAGTTGTT
TAATAGATTAGGTATTTTAAAAACTGGT**G**CATTTTTAAGTTGCTTTAAGTAAG
TTACTTAAAAGACAACAGCAGCAAAA**K**AATTAAAAAAAAATGAAAGGTGAA
GAAACACATACAAGAGAACCTTAGAACAGTAAGGTTCTAGCTAACAGGAGA
AATAAATTACAGACTGTAAAAGTTGATGACCAAGAATTTTttcagaagtggtaaaagctg
aatt
For: 5'-3' = atgctataataactaggtgttgaag
Rev 5'-3' = aattcagcttttaccacttctgaa

M144 = B9.99 (452 bp) **T to C** at position 342

AgcacaagggtcacattgagAGGTTTTAACTATAATTAAATTTTCATCTAATAAATATGA
TAATTATAAAGAAAACCAGCTGGTTTTTGGAAGACATCAAAGTGTTCTGTATC
AAGCAATAATCTCCATTAACCTATTCTGAATGGCAGGAGCAGTATGGACTGC
ATATTCTGAACTTTGGGAGGTAAATCTGTGTTGGAGCTGCTCACTGTCCATGG
AGGAGTGGAGCACAAAGTATCTGGGGGTGAAGGTCATGGCACCATTTTTCAG
CAGGGGGAGGAATAATTTTTGGTTTGAAATATTCAAAAAAAAATTTGAAAAA
ATTAAACTGGGTATGTGTG**Y**ATTTGACCATAGTAAAAAAATTTTAACAGACC
TTTTTTTGATTATCATTACATAATACAAATAAAATTTACTGATAATTCAAAAA
TTTGaacaacaaaaagccttgtcct
For: 5’-3’ = agcacaagggtcacattgag
Rev 5'-3' = aggacaaggctttttgttgtt

M145 = A8.05b (208 bp) **G to A** at position 166
TtcagcaagagtaagcaagaggCACTGAGCCGCTGGAGTCTGCACATTGATAAATTTACT
TACAGTCGTAAATAAATTGCATCATCTTCAGCTAGTAACACAGAGTCTAATTT
TTATAGCGGCATACTTGCCTCCACGACTTTCCT**A**GACACCAGAAAGAAAGGC
RAGAGCCAGCCTTAGCCTAATCaagaaccatgatccaaaaagg
For: 5’-3’ = ttcagcaagagtaagcaagagg
Rev 5'-3' = cctttttggatcatggttctt

M146 = G3.04d (395 bp) **A to C** at position 141; has(GTTTT)6 motif
GaatggggtgttacatggagaCTACAGGGCTGTTATATTCATAACTTTAGGCTATCATTAT
TGAGGGCTGGATGTCCCTCTGAGCCTCAGGATTC**AAA**GGATACT**GTTTTGTT**
TTGTTTTGTTTTGTTTTGTTTTTTCCC**M**CGGGTAATTAACACTG**G**GTTTTAG
GACAGTCTGGACTGGGGGTACATTAACAGTTGTACTAGAAACTTCCATGTCTC
AAACAGAGGGGTCTACTAGAGAAGCAATATGTCATGGAAGGCAGTTCTTCTC
CATATCTGTGTAAAGGCAAGTATTTGAAGCTAGGAGAACTGTTCCTTCTGGCC
TGTTGCCCTCTCACAGAGCACTTTAAAGTGAGCTGTGATGTGTAACTTggaaaaca
ggtctctcataatagg
For: 5’-3’ = gaatggggtgttacatggaga
Rev 5'-3' = cctattatgagagacctgttttcc

M147 = G3.35 (439 bp nominal) **1 bp insertion (extra T).** Associated with GTTT repeat. 3 T's to 4 T's at position 116. Locus also has T homopolymer which cause stutter bands during PCR.
GtattctggggcaattttaggGCAAAATACCTGAATAAGCTGGTGAAAGAAAAAAAAAGA
TACTATCAGATTAATATAAACTCATATAAGTGCAATTATGTTTTTTT**GTTTGT**
TTTGTTTTTTTCTTTCAGAGACAGGGTCTCCCTCTGTCACCTTGGCTGAAGTA
CAGTGACATGATCATGGATCACTGTAGCCTCGACCTCCTGGCCTTAAACAATC
CTTCTACCTTGGCCTCCAGAGTGGCTGGAACTACAACTGCACACCACCCCGTA
TGGCCAC**TTTTTTTTTTTT**CCCACTTTTGTAGCAATATGGTACCCAGGCTGGT
CTTGAACTCCTCTTGTCAAGCAATCTTCCTATCTTGGCCTCCCAAAATGCTTG
GATTACAGGTGTGAGCCACCACGCCTGGCCACAGTTAtgcttaaaataacctcttgtatcaa
For: 5’-3’ = gtattctggggcaattttagg
Rev 5'-3' = ttgatacaagaggttattttaagca

M147new = G3.35 (276 bp nominal) **1 bp insertion (extra T).** Associated with GTTT repeat. 3 T's to 4 T's at position 97.
GggcaaaatacctgaataagcTGGTGAAAGAAAAAAAAAGATACTATCAGATTAATATA
AACTCATATAAGTGCAATTATGTTTTTTT**GTTTGTTTTGTTT**TTTTCTTTCAG
AGACAGGGTCTCCCTCTGTCACCTTGGCTGAAGTACAGTGACATGATCATGG
ATCACTGTAGCCTCGACCTCCTGGCCTTAAACAATCCTTCTACCTTGGCCTCC
AGAGTGGCTGGAACTACAACTGCACACCACCCCGTATggccac**tTttttttttt**ccca

M148 = B9.67c (568 bp) **A to G** at position 314
AacagaattatcaggaaaaggtttCATAAAATAAAAATCTTTTAAACTTATGAAAGATGCT
CAATATAAAAAACTGTAAACCAGGGAAATGCAAATAAAAATTACAATGAAA
TACTACACACCTCCCAGAATGGCTAAAATGAAAACAAAACTGTCAATTCTAA
GTGTTAGTGAGGACATGTGGTAACCAGAACTGGCATCCAATACTAGCTGATA
AACTCGTCAATCATTTGTAAAAACAGTCTGACAATAATCCACTAGTGAAAAT
ATACATAGTCTCAGTCACAGCAATTCTATCCTGTCTATCTAGGTA**R**CAGAAAT
GTCTACATACGTTACCTAGAAACATATACTTTAATATCCACAGAATTACTTGA
AATAGCCAAAAATTGGTAACTACCAAAAGTTGAATGGTAAAACAGATAGAA
AAAAAGCTATG**C**CTAACAAAACTACACTTAATAGAACACAAGC**G**TGAGCATT
AATA**G**AACCATATAAATGCATTTTTTGAACCACTAAAAGAAGAAGCCAATAC
AAAAGAGGTGATTAAttgaaagtacacgaacaagtaaaa
For: 5'-3' = aacagaattatcaggaaaaggttt
Rev 5'-3' = ttttacttgttcgtgtactttcaa

M149 = B9.67d (568 bp) **G to A** at position 469
AacagaattatcaggaaaaggtttCATAAAATAAAAATCTTTTAAACTTATGAAAGATGCT
CAATATAAAAAACTGTAAACCAGGGAAATGCAAATAAAAATTACAATGAAA
TACTACACACCTCCCAGAATGGCTAAAATGAAAACAAAACTGTCAATTCTAA
GTGTTAGTGAGGACATGTGGTAACCAGAACTGGCATCCAATACTAGCTGATA
AACTCGTCAATCATTTGTAAAAACAGTCTGACAATAATCCACTAGTGAAAAT
ATACATAGTCTCAGTCACAGCAATTCTATCCTGTCTATCTAGGTA**A**CAGAAAT
GTCTACATACGTTACCTAGAAACATATACTTTAATATCCACAGAATTACTTGA
AATAGCCAAAAATTGGTAACTACCAAAAGTTGAATGGTAAAACAGATAGAA
AAAAAGCTATG**C**CTAACAAAACTACACTTAATAGAACACAAGC**R**TGAGCAT
TAATA**G**AACCATATAAATGCATTTTTTGAACCACTAAAAGAAGAAGCCAATA
CAAAAGAGGTGATTAAttgaaagtacacgaacaagtaaaa
For: 5'-3' = aacagaattatcaggaaaaggttt
Rev 5'-3' = ttttacttgttcgtgtactttcaa

M150 = B9.18 (289 bp) **C to T** at position 146
GcagtggagatgaagtgagacTGGGCTTTGGAGAGGTGAGGAGATGGGGCACTGACACA
CACTGCCCATGGAACCAGTCCTGACACAGGTCACACTGCAGAACTCCCACCC
CAGCTGGCACCTGCCCACACACACAGATAGAAGT**Y**GGAGAAGAGGCCATGA
GGGATGGTGCCAGTGGACTGGGCTTGGCTGAGTTGGTGCGACGCAGCTGCAG
GATACCCTCCTTCTCCTTCTGTTCCCCTTCCTTGAAGGCCACAATCTGCCATAT
Ccagaagaggggaaagtagg
For: 5'-3' = gcagtggagatgaagtgagac

Rev 5'-3' = cctactttccccctcttctg

M151 = B9.58b (422bp) **G to A** at position 209.
Acttaatttatagtttcaatccctca**G**TAATTTTAACTTACTTCTATTTTAAGAACTATAACCA
AACTATCTGTAAGACTTTTAAGCACTATCATACTCAGCTACACATCTCTTAAC
AAAAGAGGTAAATTTTGTCCTTTTTTGAA**C**GTCATAGAGTATACTCACACAAA
CCAAGAAGAAACAATCTACTACATACCTACGCTATATG**R**TATATAACTATTG
CTCCTAGGCTACAAATTAGTGCGACACTA**T**TGTACTGAATATTATAGGCCATG
TAACACAATGGTTTAAGTATCTGTGCCTCTAAACACAGAAAAGATATAGTGA
AAGTACAGTATTGCTCCTTTATTAAACTCAAAATGTTATGCAGCATATGACCG
ACTATAAAATAGCGCTTATccagatacagacatctccatgaa
For: 5'-3' = acttaatttatagtttcaatccctca
Rev 5'-3' = ttcatggagatgtctgtatctgg

M152 = B9.13 (287 bp) **C to T** at position 101
Aagctattttggtttcctttca**A**GAAAGGGCTGTGGTCTGTGGAAGGTGTCAGGAACATATT
TTCCACGGTCTGCTTTCTCCTGATAATGTTCTTCTTCT**Y**GGCCCACCTGAGAC
ATAATCCCTGAGCTCCGAGCCCTTTTTGACTGAAGCTCCTGTTGAACAAGATT
CTCAACGTTTCTACCCTGATCCACCTTCTGCCGCCGCCGTCGCCTCTCCAGAG
CCCGGCTCCTTGTCCGACTCCCTTGATGTTCAAATTTTTCCAGCTGcaatcatacccac
acaaggc
For: 5'-3' = aagctattttggtttcctttca
Rev 5'-3' = gccttgtgtgggtatgattg

M153 = A8.28c (459 bp nominal) **T to A** at position 427 bp
Ttactgataatgccatattgttttg**G**CTTAATATCAGGCTAAGTAACCACAGTATTCTGATTT**A**
AAAAAAAACATACTAGAGAGCAAGTTTATTGACAAATCTTTAGGAACTTCAG
GTACAGCATATGATTTCTGAACTATGTGTGTAAATAAGGTTTTGTTTATTCAA
ATTTAACACAGGGTAGTCTGTGTATGCCTTCCGATTTGATAGCTCTAATAAAA
CACTTTAATAGTACCATATCAAATAAATTTTATCATCATCGATTTTCTTCTTAA
TATGAAATAACACATATTTGTGATTTTTCTAAGAGTCAAAATCTCAAAAATCA
TTTTAGGTATAAAATATACCCCGAAAGTTTTATTTTATTCCATTTTATAATTAA
TCTGACTTGGAAA**GGGG**AAAAAAGCTCAAAGGGTATGTGAACA**W**TTCATTA
AGATaggaccattggtgtctgagaa
For: 5'-3' = ttactgataatgccatattgttttg
Rev 5'-3' =ttctcagacaccaatggtcct

M154 = B9.58c (422bp) **T to C** at position 252.
Acttaatttatagtttcaatccctca**G**TAATTTTAACTTACTTCTATTTTAAGAACTATAACCA
AACTATCTGTAAGACTTTTAAGCACTATCATACTCAGCTACACATCTCTTAAC
AAAAGAGGTAAATTTTGTCCTTTTTTGAA**C**GTCATAGAGTATACTCACACAAA
CCAAGAAGAAACAATCTACTACATACCTACGCTATATG**G**TATATAACTATTG
CTCCTAGGCTACAAATTAGTGCGACACTA**Y**TGTACTGAATATTATAGGCCAT
GTAACACAATGGTTTAAGTATCTGTGCCTCTAAACACAGAAAAGATATAGTG
AAAGTACAGTATTGCTCCTTTATTAAACTCAAAATGTTATGCAGCATATGACC
GACTATAAAATAGCGCTTATccagatacagacatctccatgaa

For: 5'-3' = acttaatttatagtttcaatccctca
Rev 5'-3' = ttcatggagatgtctgtatctgg

M155 = G10.57c (327 bp) **G to A** at position 251
TctctaacttctgtgagccacTCTAGCAAATTAATTGAACCAAAGGAGGAGGTTAAGGAC
AGCATAGTTTACAAAATGAGCCCTGTTTCTGACATCTGAAGTGGGGGCAGTC
TAGTGGGCCTGACCTCTTAACTTGTAGAAACATTCTTTCTTTCTAG**A**TGACTA
GTGACCAGAATTAAATTGAATCCTAGGCCACCCATTTATTGTCTTCTGCAGAA
TTGGC**G**AGAATGGAGAGGAATCCTCACCTATC**R**GTGACCAGAGATGAAATA
TTCTGAATTGAGAGTTTAAAAGAGCACACTTAGAagagatttagagtttagtttttcc
For: 5'-3' = tctctaacttctgtgagccac
Rev 5'-3' = ggaaaaactaaactctaaatctct

M156 = A8.05c (208 bp) **A to G** at position 147. Linked to M145 derived allele.
TtcagcaagagtaagcaagaggCACTGAGCCGCTGGAGTCTGCACATTGATAAATTTACT
TACAGTCGTAAATAAATTGCATCATCTTCAGCTAGTAACACAGAGTCTAATTT
TTATAGCGGCATACTTGCCTCCACGACTTTCCT**R**GACACCAGAAAGAAAGGC
GAGAGCCAGCCTTAGCCTAATCaagaaccatgatccaaaaagg
For: 5'-3' = ttcagcaagagtaagcaagagg
Rev 5'-3' = ccttttggatcatggttctt

M157 = B9.12b (352 bp) **A to C** at position 176
GctggcaagacacttctgaGCATCGGGGTGTGGACTTTACGAACCAACCTTTTAACAGT
AACTCTAGGAGAGAGGATATCAAAAATTGGCAGTGAAAAATTATAGATA**G**G
CAAAAAGCTCCTTCTGAGGTCCAGGCCAGGAGATAGTAGGATTTAAGAAACA
AACAAACAAAAAC**M**ACCACAAATGACCTTTGGTGCCACTGTCACAACTGTT
GCTCATCAGAGTAGGAGAGTTGTAGCAAAGGCATTAAAGAAGGACAAGCAG
CTGAAGAGCCTGAATCCTTGTGTTGTAAGCTATTTTGGTTTCCTTTCAAGAAA
GGGCTGTGGTCTGTggaaggtgtcaggaacatatt
For: 5'-3' = gctggcaagacacttctga
Rev 5'-3' = aatatgttcctgacaccttcc

M158 = A8.08F-newR (211 bp nominal) **G to A** at position 77, site e
tgaaatggaaatcaataaactcagtTTCCTCAAAGTTCAAAATACATGAGACTGCCTACCCTC
CTTGGAAGGCAAG**R**TGGGGCTTTCTGAAGCAAATACCAGCTTTAAAAAAAA
A**T**GTATATATATATGAAGATATATAC**AAAAAAAAAA**TTTCCCCACAACCAGA
CAATCAGAATCATCAAACCCAgaagggttaaagaaaaagaaaagg
For: 5'-3' = tgaaatggaaatcaataaactcagt
Rev: 5'-3' = ccttttctttttctttaacccttc

M159 = G10. 83new b (190 bp) **A to C** at position 89
Attggattgatttcagccttc TTCTGGTACTTTTTAAAATCTTATTAATCATTAGGAAAAGA
AGTTTTATTATTGATGCAAGCCCTAA**M**CACTCTTT**C**GACTCCAGAGGAGAAG
CTGGCAGCTCTCTGTAAGAAATATGCTGATCTTGTGAGTATTTATTTAATGGA
gcaaggaacacagaaaataaaat

For: 5'-3' = attggattgatttcagccttc
Rev 5'-3' = attttattttctgtgttccttgc

M160 = B9.47b (361 bp) **A to C** at position 251
Cagaataataggagaatttttggt**C**AAATAAAAGGCCATATTATATTTCTTTTGATAAAAGT
ATCATGTGTTCAGTATGTTTTATTATTTGAAATAATTAACATGACAGGAATAT
ATTTGAAAAAAATTCCAAAAAAAGCTAAATATACAAACTAAGAAAATTATAT
GATTATACTTATCTGCAGTATTGTAAAACAATAGTTCCAAAAACTTCTGAATT
ACAAGTTTAATACATACAACTTCAATTTTC**M**ACTACATT**GT**GGTTAGACGTT
CAGAGGAATCACAAAGGACCTCAACATGCTAGATAAGAAAATGTATTTTTTA
AATGTTTTGGCTCAgctgcttagaaaataaggaaaat
For: 5'-3' = cagaataataggagaatttttggt
Rev 5'-3' = attttccttattttctaagcagc

M161 = A8.05d original (460 bp) **C to A** at position 111
Tcacagcagcttcagcaaa**C**ACAGATTTCTGGTGTTGGAGGACAGATTTAACTACAGAA
AATTCTGTTGGGCAATCGGAAGCCTCAATCTATACAGACTTTTAGGAGGAG**M**
CTGCCTGTTTGGTTCAAATTTAGCCAAAATA**TTTTTTTTTTTAC**CACTGATTCA
GTAAATCTCCTAACTTTGCAGGAACTGGGATCCTAAAAATTATGGAACGAAT
TGTAGAAACTCAAGCAACTTTCTCCAAAGCCTAGGGttcagcaagagtaagcaagaggCA
CTGAGCCGCTGGAGTCTGCACATTGATAAATTTACTTACAGTCGTAAATAAAT
TGCATCATCTTCAgctagtaacacagagtctaatttttatAGCGGCATACTTGCCTCCACGACT
TTCCT**A**GACACCAGAAAGAAAGGC**G**AGAGCCAGCCTTAGCCTAATCaagaaccat
gatccaaaaagg
For: 5'-3' = tcacagcagcttcagcaaa
Rev: 5'-3' = ccttttggatcatggttctt
new R 5'ataaaaattagactctgtgttactagc3'(used with F primer, just amplifies the first 2 sites including homopolymer T region.

M162 = DYS257b (288 bp) =
C/T at position 202), most men are just C at position 202
Duplicated locus. Most men have both A and G alleles at position 162, however some have only the A allele. The second site at position 202 is often just C, although sometimes both C and T alleles occur on a chromosome background that is both A and G at position 162.
Gaacttgtcgggaggcaat**G**GTGACATTCATTGTGACCTTAGCCAGAGCTCACAATCAA
CCATGGTGCACTGAGACTAGCTCATGCACATTCATCAGGCAGATTCAGGCAC
CTGGCTGTCAGAGCTGTCAGCCTTCCTCAGTAGAGGAAAATGCTACAGTC**R**G
CACTGGCCTGGTATCAGGAAAATAGATGCCTGCAAAAA**Y**CCACTGTGGGACC
CTAAAAGTCTTGACCTCAGGTCCCCTTTGTGCTGTCTCTGTTGTCAGGATccacta
aaggaggaagtgtatca
For: 5'-3' = gaacttgtcgggaggcaat
Rev 5'-3' = tgatacacttcctcctttagtgg

M163 (340 bp) G10.35b **A to C** substitution at position 168

Gcagcatataaaactttcagg**A**CCCTGAAATACAGAACTGCAAAGAAACGGCCTAAGAT
GGTTGAAT**C**CTCTTTATTTTTCTTTAATTTAGACATGTTCAAACGTTCAATGTC
TTACATACTTAGTTATGTAAGTAAGGTAGCGCTTACTTCATTATGCATTTCAA
T**M**CTCAAAAAAAATTCCTTTGTGAAATGTTGAAATATTTTTCTAATCTGTTTC
ACGAGCTTCAAAAATGAGGAAAAAAGATTCAGTTTACATTTCAGCAAAATGC
CTCTTTTTAATCGGATTTATGTTTACTTAACATTTACAGTACATTTACgcttgagcaa
agttaggtttt
For: 5'-3' = gcagcatataaaactttcagg
Rev 5'-3' = aaaacctaactttgctcaagc

M164 = G10.100b (493 bp) **T to C** at position 329
TagaagtagcagattgggagaggACATGTGTTCAAGTTGTACTACTTGTATGTCTTGTTTA
GATATTACAGTCTTTTTCTTTTATCAGAAAATAATTGAATAATGATAAAATCA
GTTGCAGATTAAGACAGATTATCTGTTGCAGTCTTCTCAAAACTTAATTTAAG
TACATTATTTTCAGCTAGCATTTCTTCCTTCACATAGAACCTCCATGTGTGGA
GGGATTTCCTAATGAGTCTATTGTATGTACAATAGCACTTAATGACATAGCTT
TTAAATAATAACAGGATTTTACCAAATGTTTAATATGTGCCAGGCATCAAGC
ACC**Y**TACACAGTT**T**AATTATTGCATAGATTTGGACAGCAACTCTGCAAGTTA
GGTATGGTCATGAACCTTTGCAGATAAGGAAACTGTGTTTCACAAGGAGAAG
AAATTGTCCTGGATCATACAATAAGCTAGGATTTGCTCCAgaccatttttttcattttatcagg
For: 5'-3' = tagaagtagcagattgggagagg
Rev 5'-3' = cctgataaaatgaaaaaaatggtc

M165 = B9.008c. (340 bp) **A to G** at position 132.
Aaagcgagagattcaatccag**G**ATGACAGAATGCGTTCACCTTTAAAGGGATTAAAAGA
AGTATAATACAGTCTGTATTATTAGATCACCCAGAGACACACAAAACAAGAA
CCGT**S**AATTGAATTAGTGGTATACTAATAGAGTGGTTTTACCTGAAATATTTA
CACATCAATCCTACTGAATTCTTACAACAAATGATTTAGATTAGCTATTGTAT
TCACCAGTTGAAAGAACAGAAAATATTGAGGGAGATAACTTGTGTCAGTGCA
ACTTAATCAGATTTAGGACACAAAAGC**A**ACTACATAATGAAAAAGAGAgctggt
gacttaacttgctaaaa
For: 5'-3' = aaagcgagagattcaatccag
Rev 5'-3' = ttttagcaagttaagtcaccagc

M166 = G3.27e (393 bp) **G to A** at position 53
tggtaaactctacttagttgcctttTGGAAATGAATAAATCAAGGTAGAAAA**R**CAATTGAGA
TACTAATTCA**T**GCTCTCAGGGGAAAATCTGAATAAAGCTATCTTTTCTAACAC
AGAGCAAGTGACTCTCAAAGTCACAGTATCTGAACTAGCATATCA**G**CATCGC
CTGAATACCTAGAAATGCAAATTCCTGGGCAACACCAGAATCTAACAAAGCA
AAAAACTATGGGGGGAACAGGGAAGT**CG**GTTTAATAATACTGAGTTTGTGCA
ACCTCAACTTTGCTTTA**T**AGGAAAGCAAAATCTCAATATGATAA**A**GTTTTCTT
CAACAAAACTCTGAGATAACTATGTTGAGGGAAAGAAGTTGATCACATgcaaga
aaatctaattcgctg
For: 5'-3' = tggtaaactctacttagttgcctttt
Rev 5'-3' = cagcgaattagattttcttgc

M168 = DFFRY Ex01B site a(473 bp) **C to T** at position 371 noncoding
AgtttgaggtagaatactgtttgctGGTCTTAAAAACTGTGGTATTTTGGTGATTCCATAAAT
TAGGTCAGATACTTCCACTGGAGGGAAACAGTT**T**AAAGGATATATGTGATAC
TATTAATAGAATGAGGAAGACACACCAGATATTTAGGAGGGAATTAGCGAGC
TTGAAACTAAGAGCTGGTTTGAATGAGACTGGGTCATAAGTGATTTCAAGTA
CCAGATTAAGGCACTGAGATTTTATTTTTAAGCACTGAAGTCAGATTTTTTCC
TTTTAAAAGAAAGGATTCATGATGAAATCTGCTTTTTGTTTTGCAGAGAGCTT
GGAGATAATTCTGGTGGCTGTGTGGAGTATGTGTTGGAGGTGAGT**Y**GCTAGC
TGAAGAATTAAAACAATAGTTTTAGCAGTTTGGGTAAGAGATGTTTACAGAA
ATGTTTTGTGGAATAAAACtgaacagtcagagacctatgagatt
For: 5'-3' = agtttgaggtagaatactgtttgct
Rev: 5'-3'= aatctcataggtctctgactgttc

M169 = DFFRY Ex01B siteb (473 bp) **T to C** at position 97 noncoding
AgtttgaggtagaatactgtttgctGGTCTTAAAAACTGTGGTATTTTGGTGATTCCATAAAT
TAGGTCAGATACTTCCACTGGAGGGAAACAGTT**Y**AAAGGATATATGTGATAC
TATTAATAGAATGAGGAAGACACACCAGATATTTAGGAGGGAATTAGCGAGC
TTGAAACTAAGAGCTGGTTTGAATGAGACTGGGTCATAAGTGATTTCAAGTA
CCAGATTAAGGCACTGAGATTTTATTTTTAAGCACTGAAGTCAGATTTTTTCC
TTTTAAAAGAAAGGATTCATGATGAAATCTGCTTTTTGTTTTGCAGAGAGCTT
GGAGATAATTCTGGTGGCTGTGTGGAGTATGTGTTGGAGGTGAGT**C**GCTAGC
TGAAGAATTAAAACAATAGTTTTAGCAGTTTGGGTAAGAGATGTTTACAGAA
ATGTTTTGTGGAATAAAACtgaacagtcagagacctatgagatt
For: 5'-3' = agtttgaggtagaatactgtttgct
Rev: 5'-3'= ccagggccccgagggactctt

M170 = DFFRY Exon08 (405 bp) **A to C** at position 327
TgcttcacacaaatgcgtttcAAATAGTAACTTTTTTTCTGAAAGGGGGGAATTAATTTTT
ATTATTAACTGTATTACAGGGTTGGCTAGTGGATCTCATCAATAAATTTGGCA
CATTAAATGGGTTCCAGATTTTGCATGATCGTTTTTTTAATGGATCAGCATTA
AATATTCAAATAATTGCAGCTCTTATTAAGTAAGTTATGTTTTCATGTTTGTTA
AATAATTTCATGTTTGTTCAAATAATTGCAGCTCTTATTAAGTTATGTTTTCAT
ATTCTGTGCATTATACAAATTACTATTTTATTTACTTAAAAATCATTGTTC**M**T
TTTTTTCAGTGTGGGTTGTGTCTCACTGTAAAATGAGGACCTGTTTTTGTGTggt
cttaaatgttgaaagtaattgg
For: 5'-3' = tgcttcacacaaatgcgttt
Rev 5'-3' = ccaattactttcaacatttaagacc-3'

M171 = DFFRY Ex01B sitec (473 bp) **G to C** at position 440 noncoding?
AgtttgaggtagaatactgtttgctGGTCTTAAAAACTGTGGTATTTTGGTGATTCCATAAAT
TAGGTCAGATACTTCCACTGGAGGGAAACAGTT**T**AAAGGATATATGTGATAC
TATTAATAGAATGAGGAAGACACACCAGATATTTAGGAGGGAATTAGCGAGC
TTGAAACTAAGAGCTGGTTTGAATGAGACTGGGTCATAAGTGATTTCAAGTA
CCAGATTAAGGCACTGAGATTTTATTTTTAAGCACTGAAGTCAGATTTTTTCC
TTTTAAAAGAAAGGATTCATGATGAAATCTGCTTTTTGTTTTGCAGAGAGCTT
GGAGATAATTCTGGTGGCTGTGTGGAGTATGTGTTGGAGGTGAGT**C**GCTAGC

TGAAGAATTAAAACAATAGTTTTAGCAGTTTGGGTAAGAGATGTTTACAGAA
ATGTTTTGTG**S**AATAAAACtgaacagtcagagacctatgagatt
For: 5'-3' = agtttgaggtagaatactgtttgct
Rev: 5'-3'= ccagggccccgagggactctt

M172 = DFFRY Ex45 (345 bp) **T to G** at position197
TtgaagttacttttataatctaatgcttAATCTCTTTAAATATTTAAAATTAGGAGCCAGATGAC
CAGGATGCCCCAGATGAGCATGAGCCCTCTCCATCAGAAGATGCCCCATTAT
ATCCTCATTCACCTGCCTCTCAGTATCAACAGGTAAAAAGGATTTTTCATTTT
TATCCCCCAAACCCATTTTGATGCTT**K**ACTTAAAAGGTCTTCAATTATTATTT
TCTTAAATATTTTGAAAGTCCAAACTTTCTCTGTACCTGGCTGATATTTAAAA
CTGGATAAACTGTTCCAAACCAACATGGAGTGAAGATGGATccactgtgactgtaaagt
aataaattat
For: 5'-3' = ttgaagttacttttataatctaatgctt
Rev: 5'-3'= ataatttattactttacagtcacagtgg

M173 = DBY Ex08 (417 bp) **A to C** at position 191. Non-coding (cDNA bp# 745-52)
AagaaatgttgaactgaaagttgatGCCACTTTTCAGAAAAATGGTTGTGTTTTGTACAAAT
TGAAATACATTGTTTAAAAATAAAGCACAGTACTCACTTTAGGTTTGCCATAT
AAATTTACTGTAACTTCCTAGAAAATTGGAAATAAAGTAAGAAAAATTTTCTT
ACAATTCAAGGGCATTTAGAAC**M**CTTTGTCATCTGTTAATATTCAGAAATGA
TAAGCCAGTGTTTTGTTTTCAGGATCTGGGAAAACTGCAGCATTTCTTTTACC
CATACTGAGTCAGATATATACAGATGGTCCAGGAGAAGCTTTGAAGGCTGTG
AAGGTAAAGGTTTTGTTATAAAATCAGACATTTTTGTTTTAAAAAGCTTTGCA
AAGCCCTGTTGACTTTTCtaacggatgccagatacacct
For: 5'-3' = aagaaatgttgaactgaaagttgat
Rev: 5'-3'= aggtgtatctggcatccgtta

M174 = DffryEx38 (348 bp) **T to C** at position 219
AcatctcagatcgttgtttggtTCATAAAAATCTGTTTCTTCCATGTACCAAGCAAAATAAA
CACATCACTAAAATTTGACGTTCATAGATGTTTCTGTTTTAGGTATGATGCAC
TGTGCGTTCTTCTCCGTCACAGCAAAAATGTACGTTTTTGGTTTACTCATAAT
GTCCTTTTTAATGTATCAAATCGCTTCTCTGAATACCTTCTGGAGTGCCC**Y**AG
TGCAGAAGTGAGGGGTGCATTTGCAAAACTTATAGTGTTTATTGCACACTTTT
CCTTGCAAGATGGGTCTTGTCCTTCTCCTTTTGCATCTCCAGGACCTTCTAGTc
aggtaattgcatggcttttt
For: 5'-3' = acatctcagatcgttgtttggt
Rev: 5'-3'= aaaaagccatgcaattacctg

M175 = UTY1 exon 07 (444 bp) **5 bp deletion** at interval 84-88 non coding
TtgagcaagaaaaatagtacccaAATCAACTCAACTCCAGTGATTTAAACTCTCTGAATCA
GGCACATGCCTTCTCACTTCTC**TTCTC**AAGAATGAACAGAAACAAAGGTAT
CAGTAGAAAAAAAggtatcattaatattctttactcAAAAGTATTTCATTTAAAAATACTTAC
TTTCAGCATTGGACAAAGTACATGGATTACAGTCAATCAAGGCTAACTGAAA
ATGCTGCAAGAGAAAAGTAAAAATATTAATGCACTAAATTAAGAGTGCATAA
AAGTACATTTTCTATTTTAGCCTTTCAATGTCTATCATAAAATAACAAAGCTA

TGCTATACACCAATGCACTACACTCGACCAAATAAAATTACTGTAATTCCAA
ATTTATTTTGAAAATGTAAGTGCTAATCAAGTTATTtccctgagatagttaagaatggag
For: 5'-3' = ttgagcaagaaaaatagtaccca
Rev: 5'-3'= ctccattcttaactatctcaggga

M178 = G10.72b (514 bp) **C to T** at position 220
TaagcctaaagagcagtcagagTAGAATGCTGAATTTTCAGAAGTTTTATATTAACATAA
TCATTCATCTTTTTTGTCCTGATAATTACTCAGGAGGAAACTGAGAGGGCATG
GTCCCTTTCTATGGATAGCAATACTCAGTGTCCCAATTTTCCTTTGGGACACT
GGGACACAGGCAGAGACTCCGAAAGTCTGCATGGATTAGTTGTTCATTCACC
AYAGCTCCTTAGTGTGCCAGGAGAACTATATATGGCCTTTGGTTTCATTCAGG
GACAGGGAAACTTGAACCCATGCCTATTCATTCTCATTAAAGTAGCAGAAGT
CATGTTAGAGACAGTATTGCTGCATTCAGTACTCCTGCCTTTAACGCTTCTGA
CGCTTCCTGAAAGCAGCCCCAGCTCTCCATATGGCAAAACAAAGGCAACCTT
ATGCAAAGCCTTCTCAGGGAACCCTCAGAAAGGTTTAAACTTAGGTTCACAG
TTTTTAGAGAATAAtgtcctcattgctccctctag
For: 5'-3' = taagcctaaagagcagtcagag
Rev 5'-3' = cagagggagcaatgaggaca

M179 = Dffry exon 07 (426 bp) **C to T** at position 316
Attatgcagaattaagatgaccag TGCAGAAAAATGGAAAGAGATTATTAATAAAAATTAA
ATGTGTTTGAAATTGCAATGTGTTCTTATTATAAACTGTATCATATCCTATCCA
TGTAACAGAGATGTATTATTAACAATACTCATCGCCTAGTGGAGCTTTGTGTG
GCCAAGTTGTCCCAAGATTGGTTTCCACTTCTAGAACTTCTCGCCATGGCCTT
AAATCCTCACTGCAAGTTTCATATCTACAATGGTACACGTCCGTGTGAATTAA
TTTCCTCAAATGCTCAGTTGCCTGAAGATGAATTATTTGCTYGTTCTTCAGAT
CCTCGATCACCAAAAGTGCGTTGGTTTGTTATTTTCAAGATTAAATATTAATT
TTTTTATTTGCATTTGCCACAGAccattagtgatgtgaacctgtct
For: 5'-3' = acactactgtgctgtaatttgtgaa
Rev 5'-3' = agacaggttcacatcactaatgg

M180 = Dffry exon 11(447 bp) **T to C** at position 402
Acactactgtgctgtaatttgtgaa TGTATACATAATTTGGACTTTTGAATTCCTACTTAATA
TTATTTAGAAGTTGGAGACATGTTTTTATTTCGCTTTTTAAAAAAATTTCTTTT
TAGTTTCAGCATTGAATTTTTGTATTACATTTAGGAATGGATACAGCAAAATA
ATATCTTATCCATAGTCTTGCAAGACAGTCTTCATCAACCACAATATGTAGAA
AAGCTAGAGAAAATTCTTCGTTTTGTGATTAAAGAAAAGGCTCTTACATTAcag
gaccttgataatatctgGGCAGCACAGGTAAGAAAGTGAGATGATAGCTATTTTCTAAG
AAAGATACCAAAAAGGAGAAAATTTTTGGTAACCCTTATATAATGGCCAGCA
ATTTAGTATTGCCYGACTTTTACTAATGCATGTGctgttcatgtagagaaatcttacca
For: 5'-3' = acactactgtgctgtaatttgtgaa
Rev 5'-3' = tggtaagatttctctacatgaacag

M180 = Dffry exon 11(232 bp) **T to C** at position 128
Caggaccttgataatatctg GGCAGCACAGGTAAGAAAGTGAGATGATAGCTATTTTCTA
AGAAAGATACCAAAAAGGAGAAAATTTTTGGTAACCCTTATATAATGGCCAG

CAATTTAGTATTGCCYGACTTTTACTAATGCATGTGctgttcatgtagagaaatcttaccaAG
AATTTTTAAACAAAAAATAACATTTTTCTGTCTTTgtatatatattcatggtagcaa
NEW F 5'-3' = caggaccttgataatatctg
NEW Rev 5'-3' = ttgctaccatgaatatatatac

M181 = Dffry exon 12 (294 bp) **T to C** at position 130
GcttttatttattctacttttgttttttTCAACAGGCAGGAAAACATGAAGCCATTGTGAAGAATG
TACATGATCTGCTAGCAAAGTTGGCTTGGGATTTTTCTCCTGGACAACTTGAT
CATCTTTTTGAYTGCTTTAAGGTAGTAGCTTGAATAGTAAAGTATTGCCAAAT
AGTAAATATTGCCAGTTAATTCTAAGTAAAGTTTAATTCGTTAGATTTCTTTT
GCTTATAGCTAGTGTGCTTAACTAACATTTTCATGGAAGAATCTCTGatgaaaaaga
attggtcattgtt
For: 5'-3' = gcttttatttattctacttttgttttt
Rev 5'-3' = aacaatgaccaattctttttcat

M182 = Dffry exon 13 (364 bp) **C to T** at position 38
TattcaaagacttaaagcagtggttaATGTAAACAAAYGTAATAAATTATGTGGTATTTATA
TCATTTAAATACTTTCTTTAGGCAAGTTGGACAAATGCAAGTAAAAAGCAAC
GTGAAAAGCTCCTTGAGTTGATACGCCGTCTTGCAGAAGATGATAAAGATGG
TGTGATGGCACACAAAGTGTTGAACCTTCTTTGGAACCTGGCTCAGAGTGAT
GATGTGCCTGTAGACATCATGGACCTTGCTCTTAGTGCCCACATAAAAATACT
AGATTATAGTTGTTCCCAGGTATGGGAGTGTTTCTTTGTTCAGTTTTCTGACTT
TCCTTCACAAGTaggataacttagttacaagatgattcc
For: 5'-3' = tattcaaagacttaaagcagtggtta
Rev 5'-3' = ggaatcatcttgtaactaagttatcct

M183 = Dffry exon 19 (427 bp) **A to C** at position 324
ActgggtaaatatgactatgattgagTTACCTTTAAATTGACATTTTACTGCTTTTTATTAGAT
TGATGTCACATTTCATTTGTAAACAACCTGGATTATCTGTATTTGTCCATTATT
TATAGGTGGTTATCCATGAAGACTTCATTCAGTCTTGCTTTGATCGTTTAAAA
GCATCATATGATACACTGTGTGTTTTTGATGGTGACAAAAACAGCATTAATTG
TGCAAGACAAGAAGCCATTCGAATGGTTAGAGTATTAACTGTTATAAAAGAG
TACATTAATGAATGTGACAGTGATTATCACAAGGAAAGAATGATTCTMCCTA
TGTCGAGGTTTGTGTGAAGTTGATCTCTAGTGTTAATTTACAATTACTTAATA
TTTTCTTAGAAATTTACTTAggaaagtaataataggttaaaaggaa
For: 5'-3' = actgggtaaatatgactatgattgag
Rev 5'-3' = ttccttttaacctattattactttcc

M184 = Dffry exon 23 (305 bp) **G to A** at position 62
CactttattttagtctgtgtctttttcCTTTGCAGATAGAACAGCTGTAGAAAAATTACGARCTG
TTTGTTTGGACCATGCAAAACTTGGAGAAGGCAAACTTAGTCCACCCCTTGAC
TCTCTTTTCTTTGGTCCTTCTGCCTCCCAAGTTCTATACCTAACAGAGGTTGGT
TTTTGCCTTTGCAAAAATGTAATTTTTATATTATACGGTAATGTGAAGAACAC
TGATAAGACTGTAAAGAAAGTTTTTTAAATAGTCGAATTTCTTAGCAATGATC
agaggagaaatagatgttactaagttt
For: 5'-3' = cactttattttagtctgtgtctttttc

Rev 5'-3' = aaacttagtaacatctatttctcctct

M185 = Dffry exon 27 (430 bp) **C to T** at position 89
GgagtacctatcactgaatgtgcTTCTTAAATCCCCCTTGGAGTATATCCCAAAGAGCCTCT
CTAGCCGCAAGTGAAGAGTCTGAGGC**Y**GCATGGTCTTTACCAAGTAGGCAAT
TGTAAATGTTAACCAGAGGGTTTGTGAATTTCTTCTTGAATATGTCTCTAGGT
AACTTGCTCCTGATTCTAATTTTGCAGACCACCAATGGAAGCAATAAGCTGG
AGGTGGAAGATGAACAAGTTTGCTGTGAAGCACTGGAAGTGATGACCTTATG
TTTTGCTTTACTTCCAACAGCGTTGGATGCACTTAGTAAAGAAAAAGCCTGGC
AGACCTTCATCATTGACTTATTATTGCACTGTCCAAGCAAGTATGTGATTTTT
ATGTGTAATTTGAAGGAAGGCTTACCTTACCgttccaagcagaaatgaatgac
For: 5'-3' = ggagtacctatcactgaatgtgc
Rev 5'-3' = gtcattcatttctgcttggaac

M186 = Dffry exon 30 site a (365 bp nominal) **-1 bp deletion** (4G's to 3 G's) at position 62 (364 bp = mutant) 325 bp w/out homopolymer
TtgcatttactgttctagagagttctCAAAAAGAAATAGGAAACCACTTGAACAGTTTGGG**G**
AAGTTGTATAGAAGATCTCATTTCCTTCCAGCTCTCTGTTCTCCTAACTCCTTG
TCCTTTTCTATCTCCATGTTGTGAGTTGGGCCTATAATATTTTTCCTTTTGCAG
GATAATGTTAAAAACACAGGTGAAACAGGTGTCGAAGAGCCAATACTGGAA
GGCCACCTTGGGGTAACAAAAGAGTTATTGGCCTTTCAAACTTCTGAGAAAA
AGTATCACTTTGGTTGTGAAAAAGGAGgtgctaatctcattaaagtaagtacTTTTTTTTTTCT
TTTTTTGAgatggagtcttgctctgtgg
For: 5'-3' = ttgcatttactgttctagagagttct
Rev 5'-3' = ccacagagcaagactccatc
newRev 5'-3'=gtacttactttaatgagattagcac Homopolymer clipped off

M187 = Dffry exon 30 site b (366) **IGNORE Homopolymer in tree** T(10 to 11 T's) 325 bp w/out homopolymer
TtgcatttactgttctagagagttctCAAAAAGAAATAGGAAACCACTTGAACAGTTTGGGGA
AGTTGTATAGAAGATCTCATTTCCTTCCAGCTCTCTGTTCTCCTAACTCCTTGT
CCTTTTCTATCTCCATGTTGTGAGTTGGGCCTATAATATTTTTCCTTTTGCAGG
ATAATGTTAAAAACACAGGTGAAACAGGTGTCGAAGAGCCAATACTGGAAG
GCCACCTTGGGGTAACAAAAGAGTTATTGGCCTTTCAAACTTCTGAGAAAAA
GTATCACTTTGGTTGTGAAAAAGGAGgtgctaatctcattaaagtaagtac**T**TTTTTTTTTTCT
TTTTTTGAgatggagtcttgctctgtgg
For: 5'-3' = ttgcatttactgttctagagagttct
Rev 5'-3' = ccacagagcaagactccatc
newRev 5'-3'=gtacttactttaatgagattagcac Homopolymer clipped off

M188 = Dffry exon 31 (401 bp) **C to T** at position 185
GtattcccttgaagaaacatattgTTCCTAACCTATATTTTCTACTAATAACATGTAATGTCT
TTTTCTAACTTACTAGGAATTAATTGATGATTTCATCTTTCCCGCATCCAAAGT
TTACCTGCAGTATTTAAGAAGTGGAGAACTACCAGCTGAGCAGGCTATTCCA
GTCTGTAGTTCACC**Y**GTTACCATCAATGCCGGTTTTGAGCTACTTGTAGCATT
AGCTATTGGCTGTGTGAGGAATCTCAAACAGATAGTAGACTGTTTGACTGAA

ATGTATTACATGGGCACAGCAATTACTAGTGAGTATTTTAAATTATAAAGCTG
TTTTGTTCATTAATAATACTTCACTGTAAAATTTTATTTGGTGTTTTAgaaaaaatta
acttgtgatggactt
For: 5'-3' = gtattccctttgaagaaacatattg
Rev 5'-3' = aagtccatcacaagttaatttttc

M189 = Dffry exon 34 (378 bp) **G to T** at position 191
ActctcagcttatgtttgtcattgTTATTTTTGTTGTTATAAAATATGGATATTCTAGGCATGT
ATTACATAACTCATTTTGTTTCCTTTCCTTCTTAGGCTTTGGGGTGAACCTGTT
AATCTCCGTGAACAACATGATGCCTTAGAGTTTTTTAATTCTTTGGTGGATAG
TTTAGATGAAGCTTTAAAA**K**CTTTAGGACACCCGGCTATACTAAGTAAAGTC
CTAGGAGGCTCCTTTGCTGATCAGAAGATCTGCCAAGGCTGCCCACATAGGT
AAGTGCTAATTATGTTTTTAATGTATACTTCGTGTTGTTTTTTTTTTAATAATA
GTGTAAATCTTTCATTAGTACTTATATaaaagcagagtgtaccaaaagc
For: 5'-3' = actctcagcttatgtttgtcattg
Rev 5'-3' = gcttttggtacactctgctttt

M190 = Dffry exon 44 (346 bp) **A to G** at position 73
CtctgtcacaagtaaggaaatgatCGTGAAATTTTTGTATTAGCATTTTAAGCTGATACTGA
AAATCATTCT**R**AATTCTAAATAGTTTTATTTTTTTCTAAAGGGTAACGGAGAT
CTTAAAAGAAAATGGACCTGGGCAGTGGAATGGCTAGGAGATGAACTTGAA
AGAAGACCATATACTGGCAATCCTCAGTATAGTTACAACAATTGGTCTCCTCC
AGTACAAAGCAATGAAACAGCAAATGGTTATTTCTTAGAAAGATCACATAGT
GCTAGGATGACACTTGCAAAAGCTTGTGAACTCTGTCCAGAAGAGGTAAAAA
AAaaaaaggctaccaatggacag
For: 5'-3' = ctctgtcacaagtaaggaaatgat
Rev 5'-3' = ctgtccattggtagccttttt

M191 = DBY exon 2 (429 bp) **T to G** at position 342. Non-coding (cDNA bp# 175+120)
TtgcatttgtcatggttggtTGACCTGGACATCTTTAAAATTTGGCAGGTAATACCAGGCC
GACATGGCAGCTAAGTTTGTGGTACAGGATAAGATTGGAATCTAGGTCTCAT
TTGTCTTTTGTGATGTTATCTGTTCTTGTGTATCAGCATGTGAGCTATTGATAT
CTCTTCTAGCTTGCTAATCTGGACCTGAACTCTGAAAAACAGAGTGGAGGAG
CAAGTACAGCGAGCAGTAAGTAAAACTTTTTTTAAAAATGGAGTGTTTATCA
GAGCTTAATGTTAATGTCTTACTGGACTTGTTAATTTTAAATTTACATTTTTTT
CTTTACAACTTGACTA**K**ATGAAAATATGAGATATTTTGGTGTGTCTGGGTAAT
AAAATACACTGTTTACCTATGTCTGCTgaaaatacaaaaaattatcctggc
For: 5'-3' = ttgcatttgtcatggttggt
Rev: 5'-3'= gccaggataatttttgtattttc

M192 = DBY STS 02 (457 bp) **C to T** at position 202.
CatgggctgctgacattttgCAGGCAGGGCTCAGGGTGTTAGATGTCCTGTAATTCAGGG
ACATTCACAGTAGAAAATACTTTGGTTAGGATTTAAACCTACAAAATTGCTTT
AAACATAAACTCAAAAGTATTCTTAGGCTGGTTGCAGTGGCTTGTGTCTGCAA
TCCCAGCACTTTGGGAGGCCAAAGCAGGCAGATCC**Y**TTGAGCTCAGGAGTTT

GAGCCCAGCTTGGGCAAAATGACAAAACCCCTTCTCAGTTAAAAAAAAAAAA
TTAGCCTGGCATGGTGGGTGGTGTGCAACTGCGGTCCCAGCTACCGGGAGGC
TAAGGTGAATTACCTGAACCTGGGAGGTGGATGCTGCAGTGAGCCAAGATCC
CACCACTGCACTCCAGCCTGGATGAGGAAGTGAGATCTTGTCACAAAAACAA
AAACAAACaaacaaacaaaccaaaaggattt
For: 5'-3' = catgggctgctgacatttt
Rev: 5'-3'= aaatccttttggtttgtttgttt

M193 = DBY STS 03a (426 bp nominal) + **4 bp insertion** (CAAA) at position 56.
GcctggatgaggaagtgagTCCTGTCACAAAAACAAAAACAAACAAACAAACAAA**CA**
AACCAAAAGGATTTTTGAATACTTTAAACATACAGGGAGTGTTTTTTTTCCCC
CCGAGAAGGCAACGACTGTATAAATTTATATTGTTTTTACCATTTTAGAAATA
CTACCGTTTGCAACCCTGTTCATAATACAGTGAGTTGTGAATACATTCTGTTT
GTATTTGCAGCTAAATTAGGCAACCACTTGTGTATTTGTCAGTGTAGCAGTGG
CGGTCATTTACATGCCAAAATACATATTTTATTATAAATATTCTTTTAATTATA
TAATAATTAGGTTTGTTAGGGGCCAGAGGGGTGTCATTGTGCATCATTTGAGT
TTATTTCTTTGGGAGGCAAAGAGAGAGGAAAGGAaggtcaaaaatggagaaggc
For: 5'-3' = gcctggatgaggaagtgag
Rev: 5'-3'= gccttctccatttttgacct

M194 = DBY STS 03b (426 bp nominal) **T to C** at position 101.
GcctggatgaggaagtgagTCCTGTCACAAAAACAAAAACAAACAAACAAACAAACCA
AAAGGATTTTTGAATACTTTAAACATACAGGGAGTGTTTTTT**Y**TTCCCCCCGAG
AAGGCAACGACTGTATAAATTTATATTGTTTTTACCATTTTAGAAATACTACC
GTTTGCAACCCTGTTCATAATACAGTGAGTTGTGAATACATTCTGTTTGTATTT
GCAGCTAAATTAGGCAACCACTTGTGTATTTGTCAGTGTAGCAGTGGCGGTC
ATTTACATGCCAAAATACATATTTTATTATAAATATTCTTTTAATTATATAATA
ATTAGGTTTGTTAGGGGCCAGAGGGGTGTCATTGTGCATCATTTGAGTTTATT
TCTTTGGGAGGCAAAGAGAGAGGAAAGGAaggtcaaaaatggagaaggc
For: 5'-3' = gcctggatgaggaagtgag
Rev: 5'-3'= gccttctccatttttgacct

M195 = DBY STS 06 (515 bp nominal) **A to G** at position 430
ccactcagctttcctcaggtGCAGTCAGGTCCATCCTGCAGAGGGACCTTCTGCGGACCT
GTTCTTTCACCTCCCTAACCTGAAGATTGTATTCAAACCACCGTGGATCGCTC
ACGTAAAATGGTCACTGCGCCTAACACCTGGGATCCCGTAACCCTTATCTATC
TTGGCTTCAGAGAGTTTTTTGACTAGTTCCAACTTTGCTGAAGCTTGTCAAAG
GTAGGTGACGGCTAGTTGGAACGGAAAAATTTTACGAAACTTCCTATTCTCA
GAAGTAAAAGGGAAGAGAGAGTGCTTAAGGAAGAAGGGAAGTTGAGGGTGG
GTAAGGAGGGAGCGGGAGTTAGTGGTAGATTGTCACTGTGTTTAAGATTTCC
CCAAGGCGAAAAAGGCGAAAGATATCTTGCTAGATCCCTAGAATTCGAAGGC
ATT**R**GGAGAGGGCGGGGATAGCAAACATCGCGCGAATTTTGAGAGGCGCTG
GGACTACGTAATCCCGcgatcttatgactaaacgaacg
For: 5'-3' = ccactcagctttcctcaggt
Rev: 5'-3'= cgttcgtttagtcataagatcg

M196 = DBY STS 07 (445 bp) **C to G** at position 330.
TtagacaacttactactttgatgtcctGTTGGCTCAGTAATGCTCACGATACCAATTGTTTTGA
CAAAATAAATTTACTAAACTTGGCCTAAAATCAAACCTTGGCACAGAGGTAT
GATACAACTTTAACAGGAGTCATCAATTCATCCATAAATATAAAAAGGGAAA
AAAACTTAAGGCAGTAGTCTGCATTAGGACTGTTTGAGTTTTGCAGACTTGGG
GTTGGGAGAACATCTTAAAGCATTAAAGCATAGTTTTTTGTATGGCCAACCTT
ACTAAATTAAGTTCTGACTTGCTCACTCTATCCTGGATAGGCACTTGGGAACT
TA**S**ACTCTTTAAGCCATTCCAGTCATGATGAGGTGGAATGTATCAGTATACCA
ATTAATATTTTTGAAAGAGCTCTTTTAGGTTAATTTAAGTacagcaatttctcatgtaatgttt
a
For: 5'-3' = ttagacaacttactactttgatgtcct
Rev: 5'-3'= taaacattacatgagaaattgctgt

M197 = DBY exon 07 (408 bp) **T to C** at position 105. Non-coding (cDNA bp# 609-32)
TcagacagtttagttggttacttccATTAATATGTTAGTATAAAACAGAAATTGCGACAGAT
ACAGCATTTTATATCTGCTATGTTTACTTCTGTATTTACTTG**Y**ATTTGATTAAC
CTGGTTAAATTTCTTGGCAGTTTAGCGATATTGACATGGGAGAAATTATCATG
GGGAACATTGAACTTACTCGCTATACTCGTCCTACTCCAGTGCAAAAACATGC
CATTCCTATTATTAAGGGAAAAAGAGACTTAATGGCTTGTGCCCAAACAGGT
AAGCTTACTCAATACAAAGTGAAAGTTAAGAATACCTGATCAGACTTACTTT
AAAAGTAGTATGTTCTGAAGGGGATGTCTGAATCCTGTGTTTAGCATTTGAGG
TAGGTaaagattagctgaggatgtgtctt
For: 5'-3' = tcagacagtttagttggttacttcc
Rev: 5'-3'= aagacacatcctcagctaatcttt

M198 = DBY STS 08a (444 bp) **C to T** at position 45
TgaggtggaatgtatcagtataccAATTAATATTTTTGAAAGAG**Y**TCTTTTAGGTTAATTTA
AGTACAGCAATTTCTCATGTAATGTTTAGGGAGTTTATTCTAACCTAGGCAAA
CGGCATGCTATCACAAGAAAGGTTTAAAGCTTTGATAAAATGGGGGAGATTT
AATCAGTTTTTTTAATGCCTGCTATAAAAATTTGAAATATTAGAATGGCCGAC
CATGGCAGTGACCAGGCCTCACTACAGGCCTGGTTGGATTCTGGTCTTTAATG
CATGCTAGTGTTGATGTTTTTTGGTCAAGAACGGTTTAAACAGGAAGGATTGT
GCAGCAGGCTTTAATTTAATGTAGATTCATACTGCTCTGTTAAAGCTGCATTG
AAATGTTAAAATGGCTTACACTTGCAGACTTTGCAAATCTTaagactaacaaatccttgaa
atca
For: 5'-3' = tgaggtggaatgtatcagtatacc
Rev: 5'-3'= tgatttcaaggatttgttagtctt

M199 = DBY STS 08b (444 bp nominal) + **1 bp** insertion (extra G) at position 404 (445 bp with mutation).
TgaggtggaatgtatcagtataccAATTAATATTTTTGAAAGAGCTCTTTTAGGTTAATTTA
AGTACAGCAATTTCTCATGTAATGTTTAGGGAGTTTATTCTAACCTAGGCAAA
CGGCATGCTATCACAAGAAAGGTTTAAAGCTTTGATAAAATGGGGGAGATTT
AATCAGTTTTTTTAATGCCTGCTATAAAAATTTGAAATATTAGAATGGCCGAC
CATGGCAGTGACCAGGCCTCACTACAGGCCTGGTTGGATTCTGGTCTTTAATG
CATGCTAGTGTTGATGTTTTTTGGTCAAGAACGGTTTAAACAGGAAGGATTGT

GCAGCAGGCTTTAATTTAATGTAGATTCATACTGCTCTGTTAAAGCTGCATTG
AAATGTTAAAATGGCTTACACTTG**G**CAGACTTTGCAAATCTTaagactaacaaatcctt
gaaatca
For: 5'-3' = tgaggtggaatgtatcagtatacc
Rev: 5'-3'= tgatttcaaggatttgttagtctt

M200 = DBY STS 09a (429 bp) **G to A** at position 318
Ggcttacacttgcagactttg**C**AAATCTTAAGACTAACAAATCCTTGAAATCACACAGCTT
GCAAATACGTACTAAACTGCACAAGGTGTGTGTTCTATATGTGCAGTTTTAGC
GTATTTTAGTTGCATAGGTTTCCATGGTATTTATAGTCTCTTGTGCTAAATTTG
GCCAAAGATGATTGTCCACCACTAAAAATGCCTCTCCCACTTGGAATTCTGTA
CTGATTTTGTGGCCAGATGCAATGATCTTTAAAAACAAATCTTTTCAATGGCA
TAAGAAGTTGACAAAAATTTCTTAAAGTGCAATAGATTTTCAA**R**TGTATTGT
GCCTTGTTCTAAAACTTTTAAGTAGGTGCACTTGACAGTATTGAGGTCATTTG
TTAAGGTGCTATTTCAATTAGTGTAggtttagactcttgtacatttctcc
For: 5'-3' = ggcttacacttgcagactttg
Rev: 5'-3'= ggagaaatgtacaagagtctaaacc

M201 (326 bp) DBY exon 11&12 **G to T** at position 136
Tatgcatttgttgagtatatgtc**A**AATTGTGACACTGCAATAGTTACTACTTGAGTTACTATA
TTAGTGCAATTAATTACACAACTATATATAGTAAttagtttctcagatctaataatccagTATC
AACTGAGG**K**TTTTCGTAATAGGTACTTAGTGTTGGATGAAGCTGATAGGATG
CTGGATATGGGATTTGAACCTCAGATACGTCGTATAGTTGAACAAGATACTA
TGCCACCAAAGGGCGTTCGTCACACCATGATGTTTAGTGCTACTTTTCCTAAG
GAAATACAGGTACTGTTTGAcgtttgaactttcattcagaac
For: 5'-3' = ttagtttctcagatctaataatccagt
Rev: 5'-3'= gttctgaatgaaagttcaaacg

M202 = DBY exon 16 (392 bp) **T to G** at position 259. Non-coding (cDNA bp# 1974+38)
Ggaattgcagggtttaagc**A**GTAATTTTCAGTTTAATTGAACTTTGTACTTAACACTGCC
ATGCCATATTTTTGCTTACAGTAATAGATTCAGTGGAGGATTTGGTGCCAGAG
ACTATCGACAAAGTAGTGGTTCCAGCAGTTCTGGCTTTGGTGCTAGTCGCGGA
AGCAGCAGCCGCAGTGGTGGAGGTGGTTACGGCAACAGCAGAGGATTTGGT
GGAGGTAATGTTAATTTTTCTTTTAGGAAGGGCTTTTTGTT**K**TTCTTTTTTTTT
TTTTTTTGAGATGGAGTCCCACTCTGTCACTCAAGCTGGAGTGCAGTGGCCTG
ATCTCGGCTCACTGGAAGTGACTCTCCTGCCTCAGCCTCCTAAGTAGGTGgatt
acaggtgggtggc
For: 5'-3' = ggaattgcagggtttaagc
Rev: 5'-3'= gccacccacctgtaatcc

M203 = UTY1 exon01 (1014) (503 bp) **G to C** at position 248; synonymous substitution, SER
Gagtgccaagctgaggatga**C**CCCGTCATCAACGTGGGCAAGCTGCGTCCAGGCCTTCC
CGGAGAGTATCGCCAGCCAACCAGGCGGGTGATGGAGGTGCGTACCTGTCCA
TGCCACCAAGCGCCTCCCTTTCCTCGACTGTCAGGCTAACAGACTCCTCTTCA

CTCTCGCGGCTCGCTTTTCCTTCCGCCATTTTCTTTGCCTCATCACCGAAGGCA
ACAGCGGCGGTAGTGAGCGACACTGCGCA**S**GATTTCATGGAAACAACAAATT
TCCAAGTCCCACGACGATACCCAACCTTAATCGAGTAGTTGAAAAGACGCCT
TCAATCGCTGCTTGAGACTGTGACGCCAATTTTATCGCCTCCTCAGCGGCTGC
AAGGAAAAAAGCTGAGGCAAAGACTTAAGCTACCGAAGCACGGGCAGCGGA
ACTCGGCTACCTGGATCACATCTGGGAAACTACAGGGAAGGCAGAAGCTCGC
AGTGCtggagagcacagcagaattt
For: 5'-3' = gagtgccaagctgaggatga
Rev 5'-3': aaattctgctgtgctctcca
New Rev 5'-3':tccttggcagccgctgaggag

M204 = UTY1 ex 02 = Intron 1 (1158-4) (286 bp) **T to G** at position 234; non coding
Aaggggcgaagtattccag**AGTACGGGGACAGCAAAGGCAAGAAACACTTTTCCGACC**
CCTTGGCCATGGAGCAGAGCCAAAATAAATACTGGCTGGGCGGTAAGGAAC
GCGGGGCCTTGGTAGAGCAAAGTGCGGACCAAAGACTTTGCGTCTGGTTGCT
TTTACCTTGCCTAGTAGGGTCTTCGTTCTGGCGCCATCTTCATGAAGCCTCAC
GAACCCGAAGAGACGGCTG**K**AGAGAGAGAGACACAGAGCTTGTTAATGGTC
TGAGAAAGCCAGTGACTTGCTCCTTCCCGAGTCCAAGAGCGACAGCGACAGA
TTGGTGAGTGCCAAGCTGAGGATGACCCCGTCATCAACGTGGGCAAGCTGCG
TCCAGGCCTTCCCGGAGAGTATCGCCAGCCAACCAGGCGGGTGATGGAGGTG
CGTACCTGTCCATGCCACCAAGCGCCTCCCTTTCCTCGACTGTcaggctaacagactcct
cttca
For: 5'-3' = aaggggcgaagtattccag
Rev 5'-3': tgaagaggagtctgttagcctg

M205 = UTY Intron 2a (1221+3624) (541 bp) **T to A** at position 78.
GtataatactgtggttggaaagcaCTAAAA**T**TTAATTTTGGCTTACAGCATTATGCCTATAA
ATAAATTTTGCCACC**W**GAGTCACAGACAAAACAGGCAAAACAATCTTATTTG
GCAATTTAAATAATATCAAATGTTCCCTAGTTATTTCAATTTGACTCTTTTAAA
AGCTAGCTAGTTAGTAATAAAAGTAGGCTGGATGCAGTGGCTCACTCCTGTA
ATCCCAGCACTTTGGGAGGCTGAGGAGAGCAGATCACCTGAGGTCAGGAGTT
CCAGACCAGCCTGGCCAACATGATGAAACCCTGTCTCTACTACAAATACAAA
AAATTAGCCAAGCATGGTGGTGGATACCTGTAATCCCAGCTACTTGGGAGGC
TGAGGCAGGAGAATCACTTGAACCCAGAACACAGAGGTTGCAGTGAGGTGA
GACCGCACTATTCCACTCCAGCCAGGGCAACAAGAGTGAAACTCCATCTCGG
GGGAAAAAAAAGTAAAGTAAACCAATACCAGAAAAGTGcccatttattatcacatagtttgg
For: 5'-3' = gtataatactgtggttggaaagca
Rev 5'-3': ccaaactatgtgataataaatggg

M206 = UTY Intron 2b (1221+3671) (541 bp) **T to G** at position 31.
GtataatactgtggttggaaagcaCTAAAA**K**TTAATTTTGGCTTACAGCATTATGCCTATAA
ATAAATTTTGCCACC**T**GAGTCACAGACAAAACAGGCAAAACAATCTTATTTG
GCAATTTAAATAATATCAAATGTTCCCTAGTTATTTCAATTTGACTCTTTTAAA
AGCTAGCTAGTTAGTAATAAAAGTAGGCTGGATGCAGTGGCTCACTCCTGTA
ATCCCAGCACTTTGGGAGGCTGAGGAGAGCAGATCACCTGAGGTCAGGAGTT
CCAGACCAGCCTGGCCAACATGATGAAACCCTGTCTCTACTACAAATACAAA

AAATTAGCCAAGCATGGTGGTGGATACCTGTAATCCCAGCTACTTGGGAGGC
TGAGGCAGGAGAATCACTTGAACCCAGAACACAGAGGTTGCAGTGAGGTGA
GACCGCACTATTCCACTCCAGCCAGGGCAACAAGAGTGAAACTCCATCTCGG
GGGAAAAAAAAAGTAAAGTAAACCAATACCAGAAAAGTGcccatttattatcacatagtttgg
For: 5'-3' = gtataatactgtggttggaaagca
Rev 5'-3': ccaaactatgtgataataaatggg

M207 = UTY1 ex03 = Intron 3a (1330+18) (423 bp) **A to G** at position 79 ; non coding
Aggaaaaatcagaagtatccctg**A**AGAAGGAAAAAACGTTACAACTATGGGGCAAATGTA
AGTCAAGCAAGAAATTTA**R**AAAGAGAATAACAATACCTTTTGAATAATCTTC
CAACAAGAGGTTGAAGTGACCTAATTGGCAAAAGAAGTCAGACTCCACTTTT
CCTTCAGCTTTTAAGATTAAAGATTCGTAGCAGCGAACAGCCTAGAAATAAA
AATTATAAACATTAAGAAAAAGGCATGTCCTTCCTGGAAGAATACATACATC
TGCACGAGATTCTTAAAGAAATCAAAGCAACCATAAATGTATGTCATTTCTTC
CATAGGCATAGGATTAAATTCGGCATTTCAGAGAGGAAATAACTTCTCTTTA
AGAATTTACTAATGAAGAAATTAGATCCcaaggattcttggtgaattttg
For: 5'-3' = aggaaaaatcagaagtatccctg
Rev 5'-3': caaaattcaccaagaatccttg

M208 = UTY1 = Intron 3b (1330+5798) (507 bp) **C to T** at position 352.
Ataaatacaaaatcacctgatggat**A**TGCAAAAATTTATCAGCTTTACAAAGACATATAATA
CCATTCTATGAGCACAAGTTTATTGCAATATTTTGTCCTTTACTGTCAACAAA
AGAACACAGCCACATGATATAGGAAAAATCTATATTCTTTACAAATTTTCCAT
GAATCTCTAGCTAAAAGATCATATGACATATATGCAACGATTTATCAGCTTTC
AGAGCTTTAATTGATATTCATTACTTGTGGGTTCTGTTATTTGACTCACGAAA
ATTTATATATACACAAAATCAATACTTAATGATGGTTTCAAAGATATTCACAG
ACCTGCTCAGGGCAGCAATAAATT**Y**GACCCACTGGATACACACTCCCAGCTA
ATGTTAGAAGCGGTGGGCCTTTCTCTGACTTCATGTGTCAAGTATTCTAAACA
AACAGGCTTTTCCTGCTGTATGCAGTGTCACATTTTTCTGATTTTTGCTCttttgtta
gtaatttcgctgtttaa
For: 5'-3' = ataaatacaaaatcacctgatggat
Rev 5'-3': ttaaacagcgaaattactaacaaaa

M209 = UTY1 = Intron 3c (1330+6211) (550 bp) **A to G** at position 471.
Cactgtcttccacaatggttg**A**ACTAGTTTACAGTTCCACCAACAGTGTATAAGTTTTCCT
ATTTCTCCATATCCTCTCCAGCACCTGTTGACATTACTAAAATAACATTCTCAT
CAAGGTCATCAGGGTCTCAGAACTGGCTACATACAACCTCCAAGAAAGTTTC
GTTCTTTCTGTTTTTGCAATGTGTTCTGCCACAAATTCATCAGTTCTCAAAGCT
AACAGAACTTTTACTAGTTGCCCAATGCATCAATTCCATAGTTCTGAGAGCAT
GGGCATGAATGTCTGAAAACCTGAGGTATGATCACTAATATGCTATTCTCTGA
ACTTCTCAATTGCATTTTCCTCCTTGAATAAATCAGACTAAATTAGTGACACC
ACAAATTGTGATCATTGAGAAATCTCTAAAGGTTTTTCAGAAGCCGAGTAGG
AAGCTATCTATGACTTTTTAAAACTCTG**A**CTGAATTCT**R**AATATATTTAATTG
GACATTACATGAAGACGTTGTGTATTTAACTTCTGAATGCAgggaagataaatacaaaat
cacct

For: 5'-3' = cactgtcttccacaatggttg
Rev 5'-3': aggtgattttgtatttatcttccc

M210 = UTY1 = Intron 3d (1330+6221) (550 bp) **A to T** at position 461.
Cactgtcttccacaatggttg**A**ACTAGTTTACAGTTCCACCAACAGTGTATAAGTTTTCCT
ATTTCTCCATATCCTCTCCAGCACCTGTTGACATTACTAAAATAACATTCTCAT
CAAGGTCATCAGGGTCTCAGAACTGGCTACATACAACCTCCAAGAAAGTTTC
GTTCTTTCTGTTTTTGCAATGTGTTCTGCCACAAATTCATCAGTTCTCAAAGCT
AACAGAACTTTTACTAGTTGCCCAATGCATCAATTCCATAGTTCTGAGAGCAT
GGGCATGAATGTCTGAAAACCTGAGGTATGATCACTAATATGCTATTCTCTGA
ACTTCTCAATTGCATTTTCCTCCTTGAATAAATCAGACTAAATTAGTGACACC
ACAAATTGTGATCATTGAGAAATCTCTAAAGGTTTTTCAGAAGCCGAGTAGG
AAGCTATCTATGACTTTTTAAAACTCTG**W**CTGAATTCT**A**AATATATTTAATTG
GACATTACATGAAGACGTTGTGTATTTAACTTCTGAATGCAgggaagataaatacaaaat
cacct
For: 5'-3' = cactgtcttccacaatggttg
Rev 5'-3': aggtgattttgtatttatcttccc

M211 = UTY1 = Intron 4a (1381+16283) **C to T** at position 381.
Caattcactatttgaggaatcca**A**GTATTCCCCCTGGGGCACAGTTTAGGTATAAACACACT
TCCACTACTAACTATCTCCAGCAGTTGCCTACCTATAAGCTCCACCTACAGGC
CTGAAGTCCAGGTCACACAGCCAGCTGCAATCACTGACAACACAAGTGCACA
AACACAGGAAGCAGAACATACTACCGATGCTAGTATCACTGCACACACTACA
CTGACCACCTAGGGGCTCAGAAACTCATTTACCCACCCAATCCACTGCTACC
ACACTGGCATCTAAGAAGTCCACCCAGAGGCCCACCACGTGGTCCACCTGGA
ATTGCCAATACAGATGCTGGCAAACAATGTCGTAGGCAAAAGGATGTTAACA
ACAAG**Y**ACACCACTGAGACCAGTGAAACCTGACTACAGGCCTAACTGGCAC
TGCAGTTTCCAGCAAATTTCTCCACAGCCTCCATTAGTAACCACATCCTAGTA
TACCAAGGAAACCACAGGTACCATTAAGGGTATATActgccaaataaatcagagacttc
For: 5'-3' = caattcactatttgaggaatcca
Rev 5'-3': gaagtctctgatttatttggcag

M212 = UTY1 ex05a (409 bp) Intron 4b (1381-22) **C to A** at position 234; non coding
Tataatcaagttaccaattactggc**C**AAGATGAAAGAATGATGGGCTGAACTTGATTAGAAA
CTGCAGTAAAATAAGTGATACTACTGGAAATGTATGGTTACAGACATTAAAA
TCACCATTTACTGGAAACAAATGGTATAAGTCAACTTACCAATGAAATGCAT
TGTAGTAGAAGTAGACCAAACCAAGGCCATATAAAAACGCAGCATTCTGTTA
ATATAAAACACAAAA**M**AACCTTTATAACAGATTTTATATCTATTACTATTAC
ATATATTAATAAGAAGTCA**T**GTAACGAGATGTTTTAAGTTCTGAATATTTTAC
CATATATTACAATATTCTTCTCTACTTTTTCTCAAGTTCTCTCCATTTTGAAAA
TTGGAATCAAtttgccattcaatgttacaaaa
For: 5'-3' = tataatcaagttaccaattactggc
Rev 5'-3': ttttgtaacattgaatggcaaa

M213 = UTY1 ex05b=Intron 4c (1381-78) **T to C** at position 290. Mimics M89 (409 bp); non coding

Tataatcaagttaccaattactggc**CAAGATGAAAGAATGATGGGCTGAACTTGATTAGAAA**
CTGCAGTAAAATAAGTGATACTACTGGAAATGTATGGTTACAGACATTAAAA
TCACCATTTACTGGAAACAAATGGTATAAGTCAACTTACCAATGAAATGCAT
TGTAGTAGAAGTAGACCAAACCAAGGCCATATAAAAACGCAGCATTCTGTTA
ATATAAAACACAAAAC**AACCTTTATAACAGATTTTATATCTATTACTATTACA**
TATATTAATAAGAAGTCAY**GTAACGAGATGTTTTAAGTTCTGAATATTTTACC**
ATATATTACAATATTCTTCTCTACTTTTTCTCAAGTTCTCTCCATTTTGAAAAT
TGGAATCAAtttgccattcaatgttacaaaa
For: 5'-3' = tataatcaagttaccaattactggc
Rev 5'-3': ttttgtaacattgaatggcaaa

M214 = UTY1 ex12 = Intron 11 (1971-60) (460 bp) **T to C** at position 404; non coding
Tattacaaaatatggaaacaaggc**AACATCAAAACACAAATAGACAAACTTGCCAGCCACC**
CTTCTCCTGCCAATTATTATAGGAATATACGTGTCATTTAAAATATACTATTT
AAAATTTTTACCTGTAGAAATTTAATTCTTGCAGCAAGCGTAGAGGTATTACT
ACAACGTTTGCTTCTAGCTGCATTTAGGTAGCATTTAATGGCATCTTGAGGTT
GATTGCAGGATTCATAGAGAGTACCTAGGTCCATCCAGGCTGCGGCATGCCC
ATGGTCCAATTGTACAGCACAAATATATGCCTGTAAAGCATCCATAGGCTGA
TTTTGCTGCTGATACAACACACTGGAAAGAAAAAGAATGCTGTCAAAAACTA
CTGGTTACTTTCGTTCGTTTATTTTTCY**GTTGTTTTCAGACAGTGTCTCACACT**
GTCTCCCAGGctggagtgaagtggcatttc
For: 5'-3' = tattacaaaatatggaaacaaggc
Rev 5'-3': gaaatgccacttcactccag

M215 = UTY1 exon 14 (2358) (386 bp) **A to G** at position 163; silent substitution, SER
Gtaaaactcagatatatacatcccatg**AAATATACACAGAAACTATAAATTAGCATTAATATC**
CTCTAAAATGATACTGTAGTAAAGAAATATTCTCAAACTGTTGGTAAATTTTA
GAGAAAATAAAAATATTATACATACTTGCTGCATTAAGACAAACTGR**CTTTC**
TAACTGTTCCAGCTGATGCTTCTGTGCTGGATTTAAATTATCTCTATTTGCTCG
CAGTTGTTCCAAGTGCTAGAAGAAAAGAGATTAATATAATCAAAGTTTAATC
TAAAATTTAAGACAATATAAGGCAACTCCTCACTAAAAAGACTACACAGAAC
CTTTGCAGGATGAAAGACAGTGATTCCTAATGAAcgttaagatagtgattcttttttttt
For: 5'-3' = gtaaaactcagatatatacatcccatg
Rev 5'-3': aaaaaaaaagaatcactatcttaacg

M216 = UTY1 intron 18 3678+537 (557 bp) **C to T** at position 54.
Ctcaaccagtttttatgaagctag**AAAAAAATTCCTTTATTAAAGAAATGTAA**Y**ATTCAACA**
GGTATACATAACTAGCAGTGTCAGAATTCAGATTTAGAACCATGTTTACTAA
AAGCTTACCCTGGAACAATTATCTTTTGCTACTCTCATATAATCCCAGTCAAT
ATTTGAGAAGGCCTTAATTTTTCTAGACAAAATCTGTTTGCATATCTGGTGGT
CAAGAACCTTTTCTGTCAAAGGCCAGATAATAAATATTTTTGGCTTTATGGGC
AACCTAGTCTCTTTAGCAAACTCTGTCAATGTACTGCAAATGCAATCATAAAG
ACAGTAACTAAATAAATAAGCATAGTTATGTTCCAATAGAATTTTATTTTCAA
AAGCAGGTTGGTGGGCAGCACTTCGAGTAAGAGCATTCATTTGTTAAGTGCC
CTGAAATATAAACATGTTCTTCTGAAATATTAAACCTTTGAGAGTAAAGTCTA

TGCTCCCTAAGGCAATCTGGCTTGATTTAAAGAATACATCGATTTTCTacaagaca
cattagttcagactctc
For: 5'-3' = ctcaaccagtttttatgaagctag
Rev 5'-3': gagagtctgaactaatgtgtcttgt

M217 = UTY1 intron 17 3678+768 (461 bp) **A to C** at position 219.
GcttatttttagtctctcttccatGACTCTTCTAATA**C**CATCGTCAATAAATTTCAACTAGGTA
AAAAATTAATATTGAACATCTGTCCAAAGAAAGGCCAGTATCTCCAAAATCC
TCTCGTACAGATCTGTTTCGAGATCATTCTAATTACTGTATCTTCATATTTTAG
GTTAAGATTCTTTAACTTGTGAAGGAGAATGAAAAAGTTGGGTGACAC**M**AA
CTCTTCAGAAGGAAAAATACATAAAAATTATTTTGATGAAAGCCACAGCAGC
TTTATCAAATGCTTACGTTGCT**A**AATAGTAAAAAAAGCCACTTAAATTCCAAT
GGAAATTTTATACCCACATGTATTTATGTAAAACTTTTAAATAACATGTATTC
ATAATCACTTTTATATCCTCAACCAGTTTTTATGAAGCTAGAAAAAAATTCCT
TTATTaaagaaatgtaacattcaacaggt
For: 5'-3' = gcttatttttagtctctcttccat
Rev :5'-3': acctgttgaatgttacatttcttt

M218 = UTY1 intron 16 3679-281+768 (482 bp) **C toT** at postion 380.
TtgtgagttttttttccatcaatcTGGCTATTAAAAATCTGCAGTGCATCCTAACCTTTGATAT
TATGTTGCTACATATTACAGTATTGTATCATTTGTCTTGTCAGGAAAGTGTGG
AGGTAATAGCTAAAAAAAAACCCTCTCTTTTAAAAATTACATTTTAAATTTGAT
TCACTTTAAAACTGTTACCTATCTCTTATACCACAGTGATTTATAAAATTCTTT
TAAATTAG**T**TGAGTTGTTCGAAAGTATTTCCCAAGCATATTTTTTGAGTTATC
TTCTATTGCTTCTTAAATGAGACAACAGGTAGAAGAGACATTTAAAGTTTAA
AATCAAACTGTTTTATAAACTATTAACAAAACTTTTAG**A**GAATAAAAACCA**Y**
AACAGGCAAACCTTAAATTTGTATTTATTGCCTCAAAGTTTCAACTGAAACGC
TTATTTTTAGTCTCTCTTCCATGActcttctaataccatcgtcaataaa
For: 5'-3' = ttgtgagttttttttccatcaatc
Rev 5'-3': tttattgacgatggtattagaagag

M219 = UTY1 intron 16 3676-294 (482 bp) **T to C** at postion 232.
TtgtgagttttttttccatcaatcTGGCTATTAAAAATCTGCAGTGCATCCTAACCTTTGATAT
TATGTTGCTACATATTACAGTATTGTATCATTTGTCTTGTCAGGAAAGTGTGG
AGGTAATAGCTAAAAAAAAACCCTCTCTTTTAAAAATTACATTTTAAATTTGAT
TCACTTTAAAACTGTTACCTATCTCTTATACCACAGTGATTTATAAAATTCTTT
TAAATTAG**Y**TGAGTTGTTCGAAAGTATTTCCCAAGCATATTTTTTGAGTTATC
TTCTATTGCTTCTTAAATGAGACAACAGGTAGAAGAGACATTTAAAGTTTAA
AATCAAACTGTTTTATAAACTATTAACAAAACTTTTAG**G**GAATAAAAACCA**C**
AACAGGCAAACCTTAAATTTGTATTTATTGCCTCAAAGTTTCAACTGAAACGC
TTATTTTTAGTCTCTCTTCCATGActcttctaataccatcgtcaataaa
For: 5'-3' = ttgtgagttttttttccatcaatc
Rev 5'-3': tttattgacgatggtattagaagag

M220 = UTY1 intron 16 3676-329 (482 bp) **A to G** at postion 367.

TtgtgagttttttccatcaatcTGGCTATTAAAAATCTGCAGTGCATCCTAACCTTTGATAT
TATGTTGCTACATATTACAGTATTGTATCATTTGTCTTGTCAGGAAAGTGTGG
AGGTAATAGCTAAAAAAAACCCTCTCTTTTAAAAATTACATTTTAAATTTGAT
TCACTTTAAAACTGTTACCTATCTCTTATACCACAGTGATTTATAAAATTCTTT
TAAATTAG**C**TGAGTTGTTCGAAAGTATTTCCCAAGCATATTTTTTGAGTTATC
TTCTATTGCTTCTTAAATGAGACAACAGGTAGAAGAGACATTTAAAGTTTAA
AATCAAACTGTTTTATAAACTATTAACAAAACTTTTAG**R**GAATAAAAACCA**C**
AACAGGCAAACCTTAAATTTGTATTTATTGCCTCAAAGTTTCAACTGAAACGC
TTATTTTTAGTCTCTCTTCCATGActcttctaataccatcgtcaataaa
For: 5'-3' = ttgtgagttttttccatcaatc
Rev 5'-3': tttattgacgatggtattagaagag

M221 = UTY1 intron 18 (3784+165) (324 bp) **G *to* A** at position 200.
GggaaatgtgaaaggaaaataTCTTGGGTACCTGAAATCACTATCCTAAAGGGAAAGGT
CAAACTGGGTACTGCTTAGGGCAAACCTGCCTCCATTCTATTCAAAGTCACTC
CTCTGTTTACTGAGCTAAATGTATATCTGTTATTATCCGTATATATCTGTATAT
GATATCTATATTATCACTTGCATCAGTGCTAAAGATGCTTGCTCATGCACAAG
AGGTATAAAATTGAGTGAGAAAGAAAGATAACACACATTAAAATAAAGACT
CAGAATGTTGGGGGAAAAAATCAGTGAgtttctgtcagtgttataaaagtttaa
For: 5'-3' = gggaaatgtgaaaggaaaata
Rev 5'-3': ttaaacttttataacactgacagaaac

M223 = A8.05e (208 bp) **C to T** at position 67.
ttcagcaagagtaagcaagaggCACTGAGCCGCTGGAGTCTGCACATTGATAAATTTACTT
ACAGT**Y**GTAAATAAATTGCATCATCTTCAGCTAGTAACACAGAGTCTAATTT
TTATAGCGGCATACTTGCCTCCACGACTTTCCT**A**GACACCAGAAAGAAAGGC
GAGAGCCAGCCTTAGCCTAATCaagaaccatgatccaaaaagg
For: 5'-3' = ttcagcaagagtaagcaagagg
Rev 5'-3' = ccttttggatcatggttctt

M224 = B9.60b (301 bp) **T to C** at position 193
CttcaggcattatttttttggtTCTCCACTACAGGAGAAATGTAAATGTGATGAGTCAGAAT
TTAGGATGGCTGTATGGGTTTCTTTGACTAATACAAGAAATCACTTTGTAATG
AATGAAATCAGTGGTTTCTGCATTACTCCGTATGTTCGACATGAACACAAATT
GATACACTTAACAAAGATACTTCTTTC**Y**GCCCTTCCAAATATTTCAAAATAAG
CTGGTCATAGTACTTGCTTTTCATAAAAAGATGGTAAGCTTCCAATATTTAGA
TTTaaggaaaggtgaaggaacactat
For: 5'-3' = cttcaggcattatttttttgg
Rev 5'-3' = atagtgttccttcacctttcctt

M225= UTY1 Exon1b, (528 bp) **G to A** at position 369. (518 C to T in cDNA utr region
AaggaaaaaagctgaggcaAAGACTTAAGCTACCGAAGCACGGGCAGCGGAACTCGGC
TACCTGGATCACATCTGGGAAACTACAGGGAAGGCAGAAGCTCGCAGTGCTG
GAGAGCACAGCAGAATTTCTTAAAATCACAAACTTTGCCAGCACCAGCACAA
AGTTGTAATTGTGTCACGGGCGAACCCCACGCAGCCGCCGCGACCTCCCCGC
TCCCAACCACTTAGTTGTAGCCAATCTAGGCGACTGATTCGTCTCACGTGATC

TTTGTTGACTTACGTCAGGCATTGCTCCACTGTACTCCTAGGCTGCTGGGACC
CCGCCCAGCCAGTTCGCCAAGGACCTAGGAACATGACAGAGGCTGACT**R**ATT
CTGACCGCTGGTTGGTTGATGGTCACGTCTATGGAGAAAAGGGTAGTCTCTG
GGATGGAACAACCTGTAGGTTGTGCTAGTTAAATGCATTAAGATAGAAAATG
GAGTGTCTGTGCTGGGTGTTTTTGCAGTTGCGatacgcttgaaggggaagag
For 5'-3'= aaggaaaaaagctgaggca
Rev 5'-3'= ctcttccccttcaagcgtat

M226 UTY Ex1c 1104 silent/glu (380 bp) **C to T** at position 158
gagtgccaagctgaggatgaCCCCGTCATCAACGTGGGCAAGCTGCGTCCAGGCCTTCCC
GGAGAGTATCGCCAGCCAACCAGGCGGGTGATGGAGGTGCGTACCTGTCCAT
GCCACCAAGCGCCTCCCTTTCCTCGACTGTCAGGCTAACAGAC**YSY**TCTTCAC
TCTCGCGGCTCGCTTTTCCTTCCGCCATTTTCTTTGCCTCATCACCGAAGGCAA
CAGCGGCGGTAGTGAGCGACACTGCGCA**S**GATTTCATGGAAACAACAAATTT
CCAAGTCCCACGACGATACCCAACCTTAATCGAGTAGTTGAAAAGACGCCTT
CAATCGCTGCTTGAGACTGTGACGCCAATTTTATCGC ctcctcagcggctgcaagga
For 5'-3'=gagtgccaagctgaggatg
Rev 5'-3'=aaattctgctgtgctctcca

M227 UTY Ex1c 1105 Glu/Gln **C to G** in at position 157
GagtgccaagctgaggatgaCCCCGTCATCAACGTGGGCAAGCTGCGTCCAGGCCTTCC
CGGAGAGTATCGCCAGCCAACCAGGCGGGTGATGGAGGTGCGTACCTGTCCA
TGCCACCAAGCGCCTCCCTTTCCTCGACTGTCAGGCTAACAGAC**YSY**TCTTCA
CTCTCGCGGCTCGCTTTTCCTTCCGCCATTTTCTTTGCCTCATCACCGAAGGCA
ACAGCGGCGGTAGTGAGCGACACTGCGCA**S**GATTTCATGGAAACAACAAATT
TCCAAGTCCCACGACGATACCCAACCTTAATCGAGTAGTTGAAAAGACGCCT
TCAATCGCTGCTTGAGACTGTGACGCCAATTTTATCGCctcctcagcggctgcaagga
For 5'-3'=gagtgccaagctgaggatg
Rev 5'-3'=aaattctgctgtgctctcca

M228 UTY Ex1c (380 bp) 1106 Glu/Gly **T to C** at position 156
GagtgccaagctgaggatgaCCCCGTCATCAACGTGGGCAAGCTGCGTCCAGGCCTTCC
CGGAGAGTATCGCCAGCCAACCAGGCGGGTGATGGAGGTGCGTACCTGTCCA
TGCCACCAAGCGCCTCCCTTTCCTCGACTGTCAGGCTAACAGAC**YSY**TCTTCA
CTCTCGCGGCTCGCTTTTCCTTCCGCCATTTTCTTTGCCTCATCACCGAAGGCA
ACAGCGGCGGTAGTGAGCGACACTGCGCA**S**GATTTCATGGAAACAACAAATT
TCCAAGTCCCACGACGATACCCAACCTTAATCGAGTAGTTGAAAAGACGCCT
TCAATCGCTGCTTGAGACTGTGACGCCAATTTTATCGCctcctcagcggctgcaagga
For 5'-3'=gagtgccaagctgaggatg
Rev 5'-3'=aaattctgctgtgctctcca

M229= UTY1 Int12, **A to C** at position 159. (1560+7060 T to G in intron6)
Group I
GgtacacacctgtagtcccaacTGCTTGGGAGTCTGAGATGGAAGGATCACTTTGGGCCAG
GAATTCCACGCGTTGTACTATGATTATGCCTGTGAATAGCCACTGCACTCAAT
CCTGGAAAACAGTGAGAGCCAGTCTCTTAAAGTATAATTTCCTT**M**AATAAAAT

ATATTTCAAAATCTCTCATTCTTATTTATGATCAAAAAATGTTATTCATCAATG
TAGACTTTGAGCTTGGTCAATACTGAGCAAATAAAGCCCTCAAATATCCTTTT
CATTTGACAGGTAACTACATGCCTACTAAGGCCACGTATTATGCATATAACAA
TAAACAAACATAATCCCTCCACGAAAAAGCTCCAGCCAGAGAGAAATATTAA
AGTAAATAATTATGCTCATCTAATCCATTCAGCAATGGCAAGAATTTCACATG
AAAGTACAAGATGTCCAGCACAGATCTAACCACCTACAAATGGATGCCTCCTT
GAGAAAATGTTATTAAGGTAGGACCTGCATGGATAAGTAAAAGttaccatgaaagagtt
ctaaaaaatg
For 5'-3'=ggtacacacctgtagtcccaac
Rev 5'-3'=catttttagaactctttcatggtaa

M230 (449 bp) UTY Ex9 intron 8 1651-143 **T to A** at position 367
Group VIII
AatgtcacatttagtcttaacccatAGACTTCTAAATGAAAACAAATGTCTAAGCAGAGGGA
AAAAAATTGAACCTCAAAAGGCAAATCTCTTCAAATTAATGTAATGTATAAT
AAAAGTTTTCATGTACCTAACTGTTGCAATACAGTTGCTTTTACTTGTGCAGG
AAGGTTTTCTGTCTGCAAAAGTTGTTCATATGCCTCCTTTGCAGAATGATACT
TCCTCTAAAGAGCAAAGGAAAAAGAATATTTAGAGAAAAATAAATATTAAA
ATAAAAATACTCTTGATTTTAACAATATATACATGGCCATACTTAACTTATAA
GTAACAAATAATAAATCAATACGTAATGATGAATATTAAAAA**W**TATAAATG
TGATAATAAAAAATAAAGTAATATTACAATATTATTAAAATAGCTAgcaatgaaga
tttacatactaataatgt
For 5'-3'=aatgtcacatttagtcttaacccat
Rev 5'-3'=acattattagtatgtaaatcttcattgc

M231 UTY Ex13 Intron 13 2283+33 **G to A** at position 110 in
Group VIII
CctattatcctggaaaatgtggGCTCGTTTTAATTATATTCATATTAATTTAGTTAATCATC
ATTCAATTAATACCTAAAAAACAACATTTACTGTTTCTACTGCTTTC**R**AATTG
GGGGAAAGATCGTCAAAGAATTCATACCTGTAATTTCTGTGGTGTCAAACAC
AACGAATAAACTTGCTGTACTGGATGATGTGAAAGACTCTGGCCACCATTCC
AGTTATCAGAACCATTCTAAGGAAAATTTAGTGTAAAAGATTAAGAATATTT
GCTTAATTTCATACACTTAGAGTTATGACTAGTGAGAAccaagtgactaggaatcggaat
For 5'-3'= cctattatcctggaaaatgtgg
Rev 5'-3'=attccgattcctagtcacttgg

M232 = UTY1 intron 17 3679-566 (461 bp) **C to T** at position 38
Group VIII
gcttatttttagtctctcttccatGACTCTTCTAATA**Y**CATCGTCAATAAATTTCAACTAGGTA
AAAAATTAATATTGAACATCTGTCCAAAGAAAGGCCAGTATCTCCAAAATCC
TCTCGTACAGATCTGTTTCGAGATCATTCTAATTACTGTATCTTCATATTTTAG
GTTAAGATTCTTTAACTTGTGAAGGAGAATGAAAAAGTTGGGTGACAC**A**AAC
TCTTCAGAAGGAAAAATACATAAAAATTATTTTGATGAAAGCCACAGCAGCT
TTATCAAATGCTTACGTTGCT**A**AATAGTAAAAAAAGCCACTTAAATTCCAATG
GAAATTTTATACCCACATGTATTTATGTAAAACTTTTAAATAACATGTATTCA

TAATCACTTTTATATCCTCAACCAGTTTTTATGAAGCTAGAAAAAAATTCCTT
TATTaaagaaatgtaacattcaacaggt
For 5'-3'=gcttatttttagtctctcttccat
Rev 5'-3'= acctgttgaatgttacatttcttt

M233 = UTY1 Exon18n, **T to C** at position150, (3784+37 A to G at intron18)
Group III
AtcacttgcatcagtgctaaagaTGCTTGCTCATGCACAAGAGGTATAAAATTGAGTGAGA
AAGAAAGATAACACACATTAAAATAAAGACTCAGAATGTTGGGGGAAAAAAT
CAGTGAGTTTCTGTCAGTGTTATAAAAGTTTAAAGA**Y**AGTAAAATATATATTC
AATCTTGGTTTTAAGCTTACCTAATTTAAGAGCTCCAGCAAGGCCACGTATTA
CTGTAACAGGGTTTTTTGGATttgtacaaaattgatgtaatggagGAAAGAAAGCATCACGTT
TATTTTCCAACTGTAAAAGCAAAATATTTTGTTAGGTCTCAGATAAATGACAA
AATATACCTCAGATTTGTGCCTTTAATAAAATGATTAAATACAATACTTCAAA
TTTGTGAGTTTTTTTCCATCAATCTGGCTATTAAAAATCTGCAGTGCATCCtaacct
ttgatattatgttgctacat
For 5'-3'=atcacttgcatcagtgctaaaga
Rev 5'-3'=atgtagcaacataatatcaaaggtta

M234= UTY1 Exon20n, **C to T** at position 253, (4049 G to A in cDNA, codon 1015, Arg/Gln)
Group III
tctccattagcaatgtgtgttttACATACTGTAATTTTGCTTACATTTTTAAAAGTTTACCGGG
CATGGTGGCTCACACCTGTAATCCCAGCACTTTGGGATGCTGAGGCAAGCAGA
CCACCTGAGGTCAGGAGTTCAAGACAAGCCTGGCCAACATGGTGAAACCCTG
TCTCTACAAAAATACAAAAATTAGTTGGGCATGATGGCAGGTGCCTGTAATTC
CAGCTATTCGGGAGGCTGAGGTGGGAGAAT**Y**GCTTGAACCCAGGAGGCGGAG
GCTGCAGTGAGCTGAGATCACACCATTGCATTCCAGCCTGGGTGAGAGAGAA
TGAGACTCTGTCTCAAAAACAATAAAAAATAATAAAATAAAATAAAAGTTTA
ATAATCTATGAGCACTTTAAAAACATACTATTAACAGTATGCACTAGACAATA
ATTATGAAAGTAATATGCACTATTAAAAAATAGCAACAATTAAAAAAGGAAG
AAAGAAAAACTTACTCTCAATGATTCCTGGaaggaggaagcctggtattg
For 5'-3'=tctccattagcaatgtgtgtttt
Rev 5'-3'=caataccaggcttcctcctt

M235 = (317 bp) DFFRY Exon4, **T to G** at position 155. (1859 in cDNA, codon 65, Asp to Glu
tagatatttttccttaatctgtggtTTAAATTTGGAATATTTAATTTTTAATTAAGACTTCATCA
CCTGATTCTTCCAATGAGAATTCCGTAGCAACTCCTCCTCCAGAGGAACAAG
GGCAAGGTGATGCCCCACCACAGCATGAAGATGAAGA**K**CCTGCATTTCCACA
TACTGAGCTGGCAAACCTGGATGACATGATCAACAGGTGCATTTGTTTGGATT
TGTTTTATTAATGGATGCAGTAAACTAGAAAAGCAAAACTACTTCCAGCATT
GCAACTAGTAGTAAATgagaaaaagaaaagagtagattgtagt
For 5'-3'= tagatatttttccttaatctgtggt
Rev 5'-3'= actacaatctactcttttcttttttctc

M237= DFFRY Exon30, (366 bp) **G to C** at position 39. (5903-132 in intron29)
Group III, 325 bp w/out homopolymer region in STS.
TtgcatttactgttctagagagttctCAAAAAGAAATA**S**GAAACCACTTGAACAGTTTGGG**G**A
AGTTGTATAGAAGATCTCATTTCCTTCCAGCTCTCTGTTCTCCTAACTCCTTGT
CCTTTTCTATCTCCATGTTGTGAGTTGGGCCTATAATATTTTTCCTTTTGCAGGA
TAATGTTAAAAACACAGGTGAAACAGGTGTCGAAGAGCCAATACTGGAAGGC
CACCTTGGGGTAACAAAAGAGTTATTGGCCTTTCAAACTTCTGAGAAAAAGTA
TCACTTTGGTTGTGAAAAAGGAGgtgctaatctcattaaagtaagtac**T**TTTTTTTTTTCTTTTT
TTGAgatggagtcttgctctgtgg
For 5'-3'=ttgcatttactgttctagagagttct
newRev 5'-3'=gtacttactttaatgagattagcac Homopolymer clipped off

M238= DFFRY Exon43, **C to G** at position 28 (8729-54 in intron42)
Group I
GtactaaatggcacataattaggaaCT**S**AAATGTTAGCTACTATTGGATATTACAAAGTTTT
ACATCTGCTTCTGTTTTAGAATTCATAATGCACTTAAAGGAATTCCAGATGAC
AGAGATGGGCTGTTCGATACAATACAGCGCTC**R**AAGAATCACTATCAAAAAC
GAGCATATCAGTGCATAAAATGTATGGTAGCTCTATTTAGCAGTTGTCCTGTT
GCTTACCAGATCTTACAGGTGAGGGTTTTTCTCTTATAAATTTGTAGAAACCT
CTGTCACAAGTAAGGAAATGATCGTGAAATTTTTGTATTAGCATTTTAAgctgata
ctgaaaatcattctaaatt
For 5'-3'=gtactaaatggcacataattaggaa
Rev 5'-3'= aatttagaatgattttcagtatcagc

M239 = DFFRY Exon43, **G to A** at position 148 (8795 in cDNA, codon 2377, silent/Ser
Group I
GtactaaatggcacataattaggaaCTSAAATGTTAGCTACTATTGGATATTACAAAGTTTTA
CATCTGCTTCTGTTTTAGAATTCATAATGCACTTAAAGGAATTCCAGATGACA
GAGATGGGCTGTTCGATACAATACAGCGCTC**R**AAGAATCACTATCAAAAACG
AGCATATCAGTGCATAAAATGTATGGTAGCTCTATTTAGCAGTTGTCCTGTTG
CTTACCAGATCTTACAGGTGAGGGTTTTTCTCTTATAAATTTGTAGAAACCTC
TGTCACAAGTAAGGAAATGATCGTGAAATTTTTGTATTAGCATTTTAAgctgatac
tgaaaatcattctaaatt
For 5'-3'=gtactaaatggcacataattaggaa
Rev 5'-3'= aatttagaatgattttcagtatcagc

M240 = DBY int2n, **C to T** at position 47, (116+613 in intron1.
CtgtggaattcttgaagacgagTGACTATAATATAGCACAACGTAA**Y**AAGTATCCTGTATC
TTGTTTCTGGTGGGGTCCCGTAGCCACGGAGCAACCGTTGCCCGGGTGCTGAG
CGTGCCGAAACTGGGCTTCCGGTATGGAAAGTTTTGTGACGCAGAAGGACCG
GAAAGGGATGGTGGGGAGGGTAGGGAAGGATGGCTGCCGCGTGCTTCTCTTG
ACCCTGTAGAAATAATGGAAATTGGACGCCCGCGGAAAGACACCTGGAAGGT
TAGAGATCCAGCATTGCGCTACACCCCTTTGTTAATTCAGTCACTGGACAGCC
GCCTAGCCGAGAGCTGTGCGGTTTTTATATGGTATTGTATCTTTACTTTAGGCG
ATACATGCAGAAGTCGTCCGGTAgaaaactaacctcgaatgttgatt
For 5'-3'=ctgtggaattcttgaagacgag

Rev 5'-3'=aatcaacattcgaggttagttttc

M241 DBY Intron 4 (intron 1) **G to A** at position 57 cDNA# 117-989
AactcttgataaaccgtgctgTCTAGTTCACTAGAATTAAGTAGTAAATTCAGATG**R**CAA
GATTTTTAAGTACAGTAGTATCTTAATTGATGATTCATGTAATGTGATAGTAT
CTTGAACTTATATATGTAAGCTTTCTACGGCATAGAAAGTTTGTGCAAAAAGG
TGACCAAGGTGCT**C**TTGGCATTGGTCTTAACGTGTTTTTTGAAAAAAATCTAT
TTTAACGTACATGGTTTTTTCCCCCACCCCCGCCACCGCTTCAGAGTTGTTCTA
GGTAAGGTATTATGCTGAAAGCCCTTAAAGCGAAATAACCTTTTTTCTAGTTT
TAAAATCCATCAGTATAAGgaggcatgaattgagattgga
5'-3' For aactcttgataaaccgtgctg
5'-3' Rev tccaatctcaattcatgcctc

M242 DBY Intron 4 (intron 1) **C to T** at position 337 cDNA# 117-866
Group X
AactcttgataaaccgtgctgTCTAGTTCACTAGAATTAAGTAGTAAATTCAGATG**G**CAA
GATTTTTAAGTACAGTAGTATCTTAATTGATGATTCATGTAATGTGATAGTAT
CTTGAACTTATATATGTAAGCTTTCTACGGCATAGAAAGTTTGTGCAAAAAGG
TGACCAAGGTGCT**Y**TTGGCATTGGTCTTAACGTGTTTTTTGAAAAAAATCTAT
TTTAACGTACATGGTTTTTTCCCCCACCCCCGCCACCGCTTCAGAGTTGTTCTA
GGTAAGGTATTATGCTGAAAGCCCTTAAAGCGAAATAACCTTTTTTCTAGTTT
TAAAATCCATCAGTATAAGgaggcatgaattgagattgga
5'-3' For aactcttgataaaccgtgctg
5'-3' Rev tccaatctcaattcatgcctc

M243= DBY int6, (401 bp) **T to C** at position 142, (117-356 in intron1)
Group III
ttttgagcttttgatgtttaggaATTTATCTGCATTAAAAATAGTTGTACCGTCTTCAGGGCAA
AGATAAATTAAGGAATCTTCAAATGATTTTAATGTCCATTTATTTTTAGGGTTA
GAATATCAAGAAAACCACTGTCA**Y**TGGGAACATTTCACTATCATGACTGTAGC
TA**A**ATTGGATGTTGAAGTTACTGAGAAATTGATGGTAAATTTTTTTAGTTAGG
AAAGTTTTCACTTCGGAAAATTGTTAAGGAAAATTTGTTTTGAATTAATGAAT
TTGAACTCATTACTGTGAAACTGCTGGTATTCAGCTGATGCCATTTGCATTTGT
CATGGTTGGTAGACCTGGACATCTTTAAAATTTGGCAGGTAATACCAGGCcgaca
tggcagctaagtttg
For 5'-3'=ttttgagcttttgatgtttagga
Rev 5'-3'=caaacttagctgccatgtcg

M244= DBY int6, (401 bp) **A to C** at position 174, (117-323 in intron1)
Group I
ttttgagcttttgatgtttaggaATTTATCTGCATTAAAAATAGTTGTACCGTCTTCAGGGCAA
AGATAAATTAAGGAATCTTCAAATGATTTTAATGTCCATTTATTTTTAGGGTTA
GAATATCAAGAAAACCACTGTCA**T**TGGGAACATTTCACTATCATGACTGTAGC
TA**M**ATTGGATGTTGAAGTTACTGAGAAATTGATGGTAAATTTTTTTAGTTAGG
AAAGTTTTCACTTCGGAAAATTGTTAAGGAAAATTTGTTTTGAATTAATGAAT
TTGAACTCATTACTGTGAAACTGCTGGTATTCAGCTGATGCCATTTGCATTTGT

CATGGTTGGTAGACCTGGACATCTTTAAAATTTGGCAGGTAATACCAGGCcgaca
tggcagctaagtttg
For 5'-3'=ttttgagcttttgatgtttagga
Rev 5'-3'=caaacttagctgccatgtcg

M245= DBY int8, **del AAACA** at position 264, (174+779 in intron2)
Group I
gacgaagaacctaacattcagtgATAAAACCAAGCTCATCTGATTTTAAGGTGATGAGTTA
GCTATATTCCTGTGAAAGGAAATTAGTTATAAAGACATTCTTTTGAAATACTT
GGTCTTGTTTGGTTTTGGAAGATTGGGTGAGGTTAGTATTTGGATAGGAGAGT
AAGGCTGGTGGTTATTCAGTAGTATCCCTGGTTTGAGTCCAGGTTTCTTACTGT
TGTTCAACAAGGAAAGTAGTTGGTATGCTTTGAAACAAAACA**AAACA**GAAC
ACTTTTAAGTT**K**TATAAATTTATTTCAAACTTTGTCGTTATATGAACATTACAG
ATATTTAAATGGTAGAGACATTTTTGGATATTTAGTTAAATCCAAAAGTAGGA
GGTTTAGTTCAAATTTGGATTTTTGAGTTAcaaaatcaggtagttaagtactgtcta
For 5'-3'=gacgaagaacctaacattcagtg
Rev 5'-3'=tagacagtacttaactacctgattttg

M246= DBY int8, **T to G** at position 284, (174+799 in intron2)
Group I
gacgaagaacctaacattcagtgATAAAACCAAGCTCATCTGATTTTAAGGTGATGAGTTA
GCTATATTCCTGTGAAAGGAAATTAGTTATAAAGACATTCTTTTGAAATACTT
GGTCTTGTTTGGTTTTGGAAGATTGGGTGAGGTTAGTATTTGGATAGGAGAGT
AAGGCTGGTGGTTATTCAGTAGTATCCCTGGTTTGAGTCCAGGTTTCTTACTGT
TGTTCAACAAGGAAAGTAGTTGGTATGCTTTGAAACAAAACA**AAACA**GAACA
CTTTTAAGTT**K**TATAAATTTATTTCAAACTTTGTCGTTATATGAACATTACAGA
TATTTAAATGGTAGAGACATTTTTGGATATTTAGTTAAATCCAAAAGTAGGAG
GTTTAGTTCAAATTTGGATTTTTGAGTTAcaaaatcaggtagttaagtactgtcta
For 5'-3'=gacgaagaacctaacattcagtg
Rev 5'-3'=tagacagtacttaactacctgattttg

M247= DBY int9n, **T to C** at position 224, (175-693 in intron2)
Group II
AtggtagagacatttttggatatttAGTTAAATCCAAAAGTAGGAGGTTTAGTTCAAATTTGG
ATTTTTGAGTTACAAAATCAGGTAGTTAAGTACTGTCTACTTCATAAGTTCTT
TTACTTCTTAATCATAGACTGGCCTGTTGATTTAACTGAAAACACTTGATTTG
TTTTCCAGATCATTTTCACTTTCCAACTTTTCATGTGTTTTTATGGTATCACTT
YAATCTACCAGTACAGAATTTTTTTTCTTTTTTTGAGACGGAGTCTCGCTCTG
TCGCCCAGGCTGGAGTGCAGTGGCGCGATCTCGGCTCACCCCAAGCTCCCCC
TCCCAGGTTCATGCCATTCTCCTGCCTCAGCCTCCTGAGTAGCTGGGACTGCA
GGTGCCGGCCACCATGCCCGGCTAATTTTTTCTATTTTTTTTTAGTAGAGACA
GGGTTTCACCTTGTTAGCCAGGATGGTCTCGATCTCCTGACCTCGTGATCTGC
CCGCCT**T**GGCCTCCCaaagtgctgggattacaggc
For 5'-3'= atggtagagacatttttggatattt
Rev 5'-3'=gcctgtaatcccagcacttt

M248= DBY int9n, **T to C** at position 494, (175-444 in intron2)
Group VI
Atggtagagacatttttggatatttt**AGTTAAATCCAAAAGTAGGAGGTTTAGTTCAAATTTGG**
ATTTTTGAGTTACAAAATCAGGTAGTTAAGTACTGTCTACTTCATAAGTTCTT
TTACTTCTTAATCATAGACTGGCCTGTTGATTTAACTGAAAACACTTGATTTG
TTTTCCAGATCATTTTCACTTTCCAACTTTTCATGTGTTTTTATGGTATCACTTT
AATCTACCAGTACAGAATTTTTTTTCTTTTTTTGAGACGGAGTCTCGCTCTGTC
GCCCAGGCTGGAGTGCAGTGGCGCGATCTCGGCTCACCCCAAGCTCCCCCTC
CCAGGTTCATGCCATTCTCCTGCCTCAGCCTCCTGAGTAGCTGGGACTGCAGG
TGCCGGCCACCATGCCCGGCTAATTTTTTCTATTTTTTTTTAGTAGAGACAGG
GTTTCACCTTGTTAGCCAGGATGGTCTCGATCTCCTGACCTCGTGATCTGCCC
GCCTY**GGCCTCCC**aaagtgctgggattacaggc
For 5'-3'= atggtagagacatttttggatattt
Rev 5'-3'=gcctgtaatcccagcacttt

M249= DBY int10, **A to G** at position 313, (175-167 in intron2)
Group II
Tttcaccttgttagccaggat**GGTCTCGATCTCCTGACCTCGTGATCTGCCCGCCTTGGCCT**
CCCAAAGTGCTGGGATTACAGGCGTGAGCCACCGTGACCAGCCCAGTACAGA
TTTTTTAAAAGCCTCTTACTGGTTAGTTAATTTAGTATAGCACATAAGAGTCT
TTTTTCCCTAGTAGGCTTTTATACTGGGGTAATTACCATGTTTAATGGTCAGTG
TTGATTCATGAAGCAGTTATTGGAAATAGATCCTTTTAAAAGATAATTGTTAG
ATAACCACTACTAGCTACTGAAATATTTGTGGTTTGCAR**TGTATTTTAGAGTA**
AGCATTTTTTCCGCTCATCTTGCAAAGTAGTTTATTGTATAAAATACAGGTTTT
AAAAGTTTGTTTTCCAGGACCTATTTTTTAATagacattttctaaaagcagtatcttg
For 5'-3'=tttcaccttgttagccaggat
Rev 5'-3'=caagatactgcttttagaaaatgtct

M250= DBY int11n, **A to G** at position 299, (223+687 in intron3)
Group III
Taacagttgttaagattaccactttt**GGCCACATCCAATAAGCTGGTGAGATTGTCTGGTTTCA**
GCCTAAACAACTTCATTTGAAAGGTGTTGCATGAAATGCCTTAAAACACTTA
GGATGGTTTACTATTAAATTTGTAATTTAGAAAAGTTTAATTGGGGTGATGTT
TTGAGTGCTGCATATACATCAAAAAAATTCTAGGAGAAGGAAAGGTCAGGAA
AAGTATTTAAAACCAAAAGGAAAGAAGGTAATGATAAAGGGGTGTGGAGTG
GGTTTGTATTTCTATGTTTAGTCTGTR**GCCTCTTTAGGTCTGTTTATCAGAAGA**
CCACTTAGCTAATGATTGTATTATTTTTTCAGAATAACTGGAGAATTG**TTATT**
CTGAAAAAATATTGCATCTGGctggaattgcatcaaaggtt
For 5'-3'=taacagttgttaagattaccactttt
Rev 5'-3'=aacctttgatgcaattccag

M251= DBY int12n,(site a) (nominal, 418 bp) **G to A** at position 279, (223+1051 in intron3. Site within STS with a 7 T homopolymer length polymorphism allele.
aaatattgcatctggctggaA**TTGCATCAAAGGTTTATTAACTGCCTTAAGGAGAGTTGGC**
AATATTTTAGTATTTGAGGGGATGGAAGAGACCTTAAACATCTAACTTCCTAA
ATCTGGGAAGTACAATCGATTTAGTACAATAGATCTAGATTTAGGAAGTACA

ATTATTCATTTGTCTAATATTGGAGATTTAAAAGCAGGGGAAAATAACTTTAT
TAACTTGTAACTTTAAACATTCATTGAAATGTTTGAATTTAGGTAAGTGTGTG
GTTGTG**R**Agtgagtttactcttgtca**ttTTTTTTTT**ATCAGTTTGTAGACATGGAAAGTAG
GCAACAATGAGGG**TTTTTTT**GTTTTAACACAAGTATACCTTATTCTTAACGAG
CATATTaagattacatagttacttttggactt
For 5'-3'=aaatattgcatctggctgga
Rev 5'-3'=aagtccaaaagtaactatgtaatctt
New Rev 5'-3'=aatgacaagagtaaactcac to exclude poly T region

M252=DBY int12n, (419 bp)**ins T** at position 354, (223+1127 in intron3. (site b)
Homopolymer 7T's to 8T's
Group VI.
AaatattgcatctggctggaATTGCATCAAAGGTTTATTAACTGCCTTAAGGAGAGTTGG
CAATATTTTAGTATTTGAGGGGATGGAAGAGA**C**CTTAAACATCTAACTTCCTA
AATCTGGGAAGTACAATCGATTTAGTACAATAGATCTAGATTTAGGAAGTAC
AATTATTCATTTGTCTAATATTGGAGATTTAAAAGCAGGGGAAAATAACTTTA
TTAACTTGTAACTTTAAACATTCATTGAAATGTTTGAATTTAGGTAAGTGTGT
GGTTGTG**A**AGTGAGTTTACTCTTGTCATTTTTTTTTTATCAGTTTGTAGACATG
GAAAGTAGGCAACAATGAGGGTTTTTTT**T**GTTTTAACACAAGTATACCTTATT
CTTAACGAGCATATTaagattacatagttacttttggactt
For 5'-3'=aaatattgcatctggctgga
Rev 5'-3'=aagtccaaaagtaactatgtaatctt

M253 = DBY int13, (400 bp nominal) **C to T** at position 283
Group VI
gcaacaatgagggttttttgTTTTAACACAAGTATACCTTATTCTTAACGAGCATATTAAG
ATTACATAGTTACTTTTGGACTTTTAGAATTTGAGGCTATTTTAGAGGTCTGGT
AGAGCAAAGTAGACAACATGGAAATTCCTTGTTTTGTATTGACTACTTCCATT
TAGCTGATCTGTTTCTTTTTGGTGTTACTAGACAAAGCTAGATTTTAAAAGATG
AATTAAGATGCTCAGCTAACTAGTCCTGTTTATAGTATTGTTGATAGATAGCA
AGTTGA**Y**TTCTCCAGGTTCTTCATTGAATGAGTCCTTGTTTACTATGATGCTTG
CTACATACAGTTGCTACATACTACTATGTATGAGTAGTTTTTGGTCATaaactgcata
gaggtggagctg
For 5'-3'=gcaacaatgagggttttttg
Rev 5'-3'=cagctccacctctatgcagttt

M254 = DBY int13, (400 bp nominal, 418 bp derived)**18bp INSERTION + 2bp substitution**, A to G and G to C at positions 339, 340
Group VIII
gcaacaatgagggttttttgTTTTAACACAAGTATACCTTATTCTTAACGAGCATATTAAG
ATTACATAGTTACTTTTGGACTTTTAGAATTTGAGGCTATTTTAGAGGTCTGG
TAGAGCAAAGTAGACAACATGGAAATTCCTTGTTTTGTATTGACTACTTCCAT
TTAGCTGATCTGTTTCTTTTTGGTGTTACTAGACAAAGCTAGATTTTAAAAGA
TGAATTAAGATGCTCAGCTAACTAGTCCTGTTTATAGTATTGTTGATAGATAG
CAAGTTGA**C**TTCTCCAGGTTCTTCATTGAATGAGTCCTTGTTTACTATGATGCT

TGCTACATAC**TACTATGTTTACTATGATRS**TTGCTACATACTACTATGTATG
AGTAGTTTTTGGTCATaaactgcatagaggtggagctg
For 5'-3'=gcaacaatgagggtttttttg
Rev 5'-3'=cagctccacctctatgcagttt

M255= DBY int14, (within derived 471 bp) **C to T** at position 107, (224-813, in intron3)
Group V
ttttttttgagacggagtcttgCTGTGTTGTCCAGGCTGGAGTACAGTGGCGCGATCTCAGC
TCACTGCAAGCTCCACCTCTTGGGTTCATGCCATTCTCCTGCCT**Y**AGGCTCCT
GAGTAGCTGGGACTACATAGGTGCCCGCCACCATGCCCAGCTAATTTTTTTGT
ATTTTTAGTAGAGACGGGGTTTCACCGTGTTAGCCAGGATGGTCTTGATCTCC
TGACCTTGTGATCTGCCTGCCTTAGCC**C**TCCCAAAGTGCTGGGATTACAGGT
GTGAGCCATCCCTGTTTTAATCCATCTGACATATTTCTTCTGATTATGTAGCTC
TCTTAGTTCAAGCTTTTCTGTAGGTAACCCACAGTCCCTGAGGTAA**T**CTTTTA
CTTAGCTGGGCCTTCCCAAAATGTGTATTATATATAGCATATGTTAAATGTTT
AGGTTTAACACCTtttgtattattcaggatttgtcaag
For 5'-3'=ttttttttgagacggagtcttg
Rev 5'-3'=cttgacaaatcctgaataatacaaa

M256 = DBY int14, (derived 471 bp) **ins C** at position 249, (224-672 in intron3)
Group V
ttttttttgagacggagtcttgCTGTGTTGTCCAGGCTGGAGTACAGTGGCGCGATCTCAGC
TCACTGCAAGCTCCACCTCTTGGGTTCATGCCATTCTCCTGCCT**C**AGGCTCCT
GAGTAGCTGGGACTACATAGGTGCCCGCCACCATGCCCAGCTAATTTTTTTGT
ATTTTTAGTAGAGACGGGGTTTCACCGTGTTAGCCAGGATGGTCTTGATCTCC
TGACCTTGTGATCTGCCTGCCTTAGCC**C**TCCCAAAGTGCTGGGATTACAGGT
GTGAGCCATCCCTGTTTTAATCCATCTGACATATTTCTTCTGATTATGTAGCTC
TCTTAGTTCAAGCTTTTCTGTAGGTAACCCACAGTCCCTGAGGTAA**T**CTTTTA
CTTAGCTGGGCCTTCCCAAAATGTGTATTATATATAGCATATGTTAAATGTTT
AGGTTTAACACCTtttgtattattcaggatttgtcaag
For 5'-3'=ttttttttgagacggagtcttg
Rev 5'-3'=cttgacaaatcctgaataatacaaa

M257= DBY int14, (nominal 470 bp) **T to C** at position 373, (224-547 in intron3)
Group I
ttttttttgagacggagtcttgCTGTGTTGTCCAGGCTGGAGTACAGTGGCGCGATCTCAGC
TCACTGCAAGCTCCACCTCTTGGGTTCATGCCATTCTCCTGCCT**C**AGGCTCCT
GAGTAGCTGGGACTACATAGGTGCCCGCCACCATGCCCAGCTAATTTTTTTGT
ATTTTTAGTAGAGACGGGGTTTCACCGTGTTAGCCAGGATGGTCTTGATCTCC
TGACCTTGTGATCTGCCTGCCTTAGCCTCCCAAAGTGCTGGGATTACAGGTGT
GAGCCATCCCTGTTTTAATCCATCTGACATATTTCTTCTGATTATGTAGCTCTC
TTAGTTCAAGCTTTTCTGTAGGTAACCCACAGTCCCTGAGGTAA**Y**CTTTTACT
TAGCTGGGCCTTCCCAAAATGTGTATTATATATAGCATATGTTAAATGTTTAG
GTTTAACACCTtttgtattattcaggatttgtcaag
For 5'-3'=ttttttttgagacggagtcttg

Rev 5'-3'=cttgacaaatcctgaataatacaaa

M258=DBY int15, (475 bp) **T to C**, at position 123, (224-388, in intron3)
Group VI
Tatatagcatatgttaaatgtttaggt**T**TAACACCTTTTGTATTATTCAGGATTTGTCAAGGATG
GGACATAACTAAGAAACTAACAATGGGCTTGCACTAGCTACAAGTTCAGCTT
AAAAA**Y**TGGGAACTTGGAATCCCTCTTAGTCATAGCTTAAAAAAAGACTCAT
CTTAAATAATTTAATTGGAGTAGGTTTATATTTTGGATATGTAACATTTACAC
TTAAAAAATGAATGAAAAAAATTGTTACGATAGTATAGTATTAATAGCATAG
CTATGTTACATGCAAGCTACCTTGTTCTCAGGTCATGAGATTACTTTGCTTCAT
ATAATAATCTCTGGTGGAAGAAAACATTAAAGCTTTTAACAATTCTGCTTATG
GGACTTGTAGACCATTGGTCCCATAAAGATAACATAAAGGAAGACTACATGT
GAAGGACTTCATATTTTgaaagatgcaaattattcaaaagtc
For 5'-3'=tatatagcatatgttaaatgtttaggt
Rev 5'-3'=gacttttgaataatttgcatctttc

M259= DBY int16, (396 bp)**T to G** at position 151, (352+271, in intron4
Group IX
Cagaatgttggtttactcattgtt**T**TGTTAGCAGTAAGAGGTCTTTATTAATTTATTAAATTA
GATGAATATGGTATTTGACACAGTGAAATCTGTTTCAACTTAAATGATACTTA
AAGCCTGTCTGTGACAGCTTTAAACACTTCATTT**K**TGATGTGTGTTATAAGTT
GATCTTAAAAACCTAATGGCTGTATTTAATCCTTTCTGTTTTTCACAAATAGG
AGTAAAACTCTAAAAATATTCTCTTGTCACATGTCTACTTTCATATAAAGGAG
AAATTCAAGTGTTATTCCTGCTTTCCTACTAGTAAATATATTTAGATGATACT
ATTTTAAATGAAGATGTAAAGTACGTAACTAGTTATAAGTATCTaaaaacctaattctt
agcatgtga
For 5'-3'=cagaatgttggtttactcattgtt
Rev 5'-3'=tcacatgctaagaattaggttttt

M260= DBY int19, (343 bp) **G to A** at position 253, (608-124 in intron6)
Group VI
Ccacacccagctcatttttt**G**TACTTTTAGTAGAGACAGGGTTTCGCCATGTTGGCCAGGC
TGGTCTCAAATTCCTGATCTCAAGTGATCTTCATGCCTTAGCCTCCCAGAGTG
CTGGGACTACAGGCATCAGCCACCATACCTGGCCTCCAAAAACTTTTTTCAAT
GTAGATTAAACCCAGGCATTTTCTTAAAAAATGCCATGAATCTTTTACTGAAA
TCATAGCATCTGTAAACTAAATCAGACAGTTTA**R**TTGGTTACTTCCATTAATA
TGTTAGTATAAAACAGAAATTGCGACAGATACAGCATTTTATATctgctatgtttacttc
tgtatttactt
For 5'-3'=ccacacccagctcatttttt
Rev 5'-3'=aagtaaatacagaagtaaacatagcag

M261= DBY int22, (284 bp) **A to G** at position 213, (1090-32 in intron10)
Group X
Atttgaggctctgagcttca**T**TTTAACAATCAACATGGGTAATTCGGTTGTTACCTTGAGC
ATTTCATCTCATGATTTTGTGTGTGTTTGTGTGTGTATGCATTTGTTGAGTATA
TGTCAAATTGTGACACTGCAATAGTTACTACTTGAGTTACTATATTAGTGCAA

TTAATTACACAACTATATATAGTAATTAGTTTCTCAGATCTAAT**R**ATCCAGTA
TCAACTGAGGGTTTTCGTAATAGGTACTTAGTGTTGGATGAAgctgataggatgctggat
atg
For 5'-3'=atttgaggctctgagcttca
Rev 5'-3'=catatccagcatcctatcagc

M262= DBY STS01, (502 bp) **del A** at position 226, (1-2908 out side of 5' region) Group III
agctgtttggacttgagtagttgTAGAATAACTGAAAATAGGAAACTGCTATATATATATGT
ATGTATAATATATATAACCTTTTTCAGGTACTCCTATTGCAATACCTGCATTT
CAGCACTATTCAAAAGTAAAATAAGTCCCAGAGCCAGGTTAGTCATTATGTC
CTATTTATTGCTAATTTCATATACAAATGAGAGCTGTCAGAATTCACAGCTT
CTGA**A**TATCAGAAGCTCATGTTTTCCCTGGTCTATACAAAAAGGAAATAAGT
GAGGCCAAAAATGTACTTTAACAGTGCTCCATAATACGAATCTCATAAATGA
GCTGGAATAGACCCTGAGGTCTTCAAGCCTAGTTTCTCAAGATCGTATTTTGT
AAACTTGTGCTAGCAGTTTTGAATATCACAATGATTGGCATGGGCTGCTGACA
TTTTAGCAGGCAGGGCTCAGGGTGTTAGATGTCCTGTAATTCAGGgacattcacagta
gaaaatactttgg
For 5'-3'= agctgtttggacttgagtagttg
Rev 5'-3'= ccaaagtattttctactgtgaatgtc

M263=DBY STS06, (515 bp) **G to C** at position 332, (1-341 out side of 5' region) Group III
ccactcagctttcctcaggtGCAGTCAGGTCCATCCTGCAGAGGGACCTTCTGCGGACCT
GTTCTTTCACCTCCCTAACCTGAAGATTGTATTCAAACCACCGTGGATCGCTC
ACGTAAAATGGTCACTGCGCCTAACACCTGGGATCCCGTAACCCTTATCTATC
TTGGCTTCAGAGAGTTTTTGACTAGTTCCAACTTTGCTGAAGCTTGTCAAAG
GTAGGTGACGGCTAGTTGGAACGGAAAAATTTTACGAAACTTCCTATTCTCA
GAAGTAAAAGGGAAGAGAGAGTGCTTAAGGAAGAAGGGAAGTTGAGGGTGG
GTAAGGAGG**S**AGCGGGAGTTAGTGGTAGATTGTCACTGTGTTTAAGATTTCC
CCAAGGCGAAAAAGGCGAAAGATATCTTGCTAGATCCCTAGAATTCGAAGGC
ATTAGGAGAGGGCGGGGATAGCAAACATCGCGCGAATTTTGAGAGGCGCTG
GGACTACGTAATCCCGcgatcttatgactaaacgaacg
For 5'-3'= ccactcagctttcctcaggt
Rev 5'-3'= cgttcgtttagtcataagatcg

M264=DBY Exon17, (552 bp) **C to T** at position 115, (1988 at cDNA, codon639, silent/Gly)
Group III.
tccaactctagatttcttttactggTTTTATGTTAAAGTACTTGAGAAAAAAAAGGTATTAAC
GAATGACTTAATTTCTCTCTAAACATTTTTCTTGATAGGTGGCTATGGAGG**Y**T
TCTACAATAGTGATGGATATGGAGGAAATTATAACTCCCAGGGGGTTGACTG
GTGGGGCAACTGAATCTGCTTTGCAGCAAAGTCACCCTTACAAAGAAGCTAA
TATGGAAACCACATGTAACTTAGCCAGACTATATTGTGTAGCTTCAAGAACTT
GCAGTACATTACCAGCTGTGATTCTCCTGATAATTCAAGGGAGCTCAAAGTC
ACAAGAAGAAAAATGAAAGGAAAAAACAGCAGCCCTATTCAGAAATTGGTT

TGAAGATGTAATTGCTCTAGTTTGGATTAAACTCTTCCCCTCCTGCTTTAGTGC
CACCCCAAACTGCATTTATAATTTTGTGACTGAGGATCGTTTGTTTGTTAACG
TACTGTGACTTTAACTTTAGACAACTTACTACTTTGATGTCCTGTTGgctcagtaatg
ctcacgatacc
For 5'-3'=tccaactctagatttcttttactgg
Rev 5'-3'=ggtatcgtgagcattactgagc

M265= DBY STS07, **C to A** at position 298, (2312+358 outside 3' region)
ttagacaacttactactttgatgtcctGTTGGCTCAGTAATGCTCACGATACCAATTGTTTTGAC
AAAATAAATTTACTAAACTTGGCCTAAAATCAAACCTTGGCACAGAGGTATG
ATACAACTTTAACAGGAGTCATCAATTCATCCATAAATATAAAAAGGGAAAA
AAACTTAAGGCAGTAGTCTGCATTAGGACTGTTTGAGTTTTGCAGACTTGGGG
TTGGGAGAACATCTTAAAGCATTAAAGCATAGTTTTTTGTATGGCCAACCTTA
CTAAATTAAGTTCTGACTTGCT**M**ACTCTATCCTGGATAGGCACTTGGGAACTT
A**C**ACTCTTTAAGCCATTCCAGTCATGATGAGGTGGAATGTATCAGTATACCA
ATTAATATTTTTGAAAGAGCTCTTTTAGGTTAATTTAAGTacagcaatttctcatgtaatgttt
a
For 5'-3'=ttagacaacttactactttgatgtcct
Rev 5'-3'=taaacattacatgagaaattgctgt

M266= DBY STS08, (444 bp) **T to C** at position 208, (2312+623 outside 3' region)
Group II
tgaggtggaatgtatcagtataccAATTAATATTTTTGAAAGAG**C**TCTTTTAGGTTAATTTAA
GTACAGCAATTTCTCATGTAATGTTTAGGGAGTTTATTCTAACCTAGGCAAAC
GGCATGCTATCACAAGAAAGGTTTAAAGCTTTGATAAAATGGGGGAGATTTA
ATCAGTTTTTTTAATGCCTGCTATAAAAATTTGAAATAT**Y**AGAATGGCCGACC
ATGGCAGTGACCAGGCCTCACTACAGGCCTGGTTGGATTCTGGTCTTTAATGC
ATGCTAGTGTTGATGTTTTTTGGTCAAGAACGGTTTAAACAGGAAGGATTGTG
CAGCAGGCTTTAATTTAATGTAGATTCATACTGCTCTGTTAAAGCTGCATTGA
AATGTTAAAATGGCTTACACTTGCAGACTTTGCAAATCTTaagactaacaaatccttgaaat
ca
For 5'-3'=tgaggtggaatgtatcagtatacc
Rev 5'-3'=tgatttcaaggatttgttagtctt

M267 EIF1A Y STS12 (site a) (287 bp) **T to G** at position 148. STS also contains two Group I associated mutations
ttatcctgagccgttgtccctgTGTTTCCATTTCTCTTTTCCTCATTTCTCATCATC**T**ACATTT
CTCCT**G**TACTTGTTCATTAAATAATGATTCCTTGG**A**TATACCAAGTCTGGATA
GCGGATTCGAT**G**GAAGCATTTTTGTAAATA**K**AC**G**TTCAGTATTTTGTGTGGA
AGAACACAATCTAGCTGATGCCTGCAATCCCAGCCCTTTGGAAAGCGAGGTG
GGTGGATTGCTTGAAGCTACGAGTTTGACACTAGCCTGGGCAACagggtacaaccgt
gtctctaca
newFor 5'-3=ttatcctgagccgttgtccctg
Rev 5'-3'=tgtagagacacggttgtaccct

M268 = EIF1A_Y STS5a, (427 bp) **A to G** at position 292,

GROUP VII
ctaaagatcagagtatctcccttgCAAAATGTCCATTAAATCTTTGCTGATGTTATTATCCCT
GTACCTGACTCTATCCTTAAATAGTAAGGCTTCCTTTATTCTTGTAGGGTAGA
ACTTTTAAACTGAGTGATGCCTAAAAATGTTCTCAATAAAGAGAGTATCTCCA
AAACACGTCGGATTTGTTTAAAGAGGAAGTGTGGATTTTTTGATCTTAGAAA
GGAAACGAGATAAAATATTAAACGACTTTAATTTTTGTATGATCATGCCTAGC
CTCATTCCTCTAAAAT**R**TAATTTAAAGTGGATTCTGTTACATGGTATCACAAT
AGAAGGGGAATGATCAGGGTTTGGTTAAT**T**CTGGTAAATTGAAAACAATTTT
TTTTTT(**T**)ATCATATGTGCCTCAgaaggcacacaaaagaagtatagt
For: 5'-3'= ctaaagatcagagtatctcccttg
Rev: 5'-3'= actatacttcttttgtgtgccttc

M269 = EIF1A_Y STS5b, (427 bp) **T to C** at position 358,
Group IX
CtaaagatcagagtatctcccttgCAAAATGTCCATTAAATCTTTGCTGATGTTATTATCCC
TGTACCTGACTCTATCCTTAAATAGTAAGGCTTCCTTTATTCTTGTAGGGTAG
AACTTTTAAACTGAGTGATGCCTAAAAATGTTCTCAATAAAGAGAGTATCTCC
AAAACACGTCGGATTTGTTTAAAGAGGAAGTGTGGATTTTTTGATCTTAGAA
AGGAAACGAGATAAAATATTAAACGACTTTAATTTTTGTATGATCATGCCTA
GCCTCATTCCTCTAAAATATAATTTAAAGTGGATTCTGTTACATGGTATCACA
ATAGAAGGGGAATGATCAGGGTTTGGTTAAT**Y**CTGGTAAATTGAAAACAATT
TTTTTTTT(**T**)ATCATATGTGCCTCAgaaggcacacaaaagaagtatagt
For: 5'-3'= ctaaagatcagagtatctcccttg
Rev: 5'-3'= actatacttcttttgtgtgccttc

M270 = EIF1A_Y STS5, (428 bp) **ins T** at position 387.. Has ancestral T at M281.
HOMOPOLYMER
CtaaagatcagagtatctcccttgCAAAATGTCCATTAAATCTTTGCTGATGTTATTATCCC
TGTACCTGACTCTATCCTTAAATAGTAAGGCTTCCTTTATTCTTGTAGGGTAG
AACTTTTAAACTGAGTGATGCCTAAAAATGTTCTCAATAAAGAGAGTATCTCC
AAAACACGTCGGATTTGTTTAAAGAGGAAGTGTGGATTTTTTGATCTTAGAA
AGGAAACGAGATAAAATATTAAACGACTTTAATTTTTGTATGATCATGCCTA
GCCTCATTCCTCTAAAATATAATTTAAAGTGGATTCTGTTACATGGTATCACA
ATAGAAGGGGAATGATCAGGGTTTGGTTAAT**T**CTGGTAAATTGAAAACAATT
TTTTTTTT**T**ATCATATGTGCCTCAgaaggcacacaaaagaagtatagt
For: 5'-3'= ctaaagatcagagtatctcccttg
Rev: 5'-3'= actatacttcttttgtgtgccttc

M271 = UTY1 intron 17 3679-566 (461 bp) **A to C** at position 296
Group VIII. Discovered while typing M232. This STS also contains M217 site.
gcttatttttagtctctcttccatGACTCTTCTAATA**C**CATCGTCAATAAATTTCAACTAGGTA
AAAAATTAATATTGAACATCTGTCCAAAGAAAGGCCAGTATCTCCAAAATCC
TCTCGTACAGATCTGTTTCGAGATCATTCTAATTACTGTATCTTCATATTTTAG
GTTAAGATTCTTTAACTTGTGAAGGAGAATGAAAAAGTTGGGTGACAC**A**AAC
TCTTCAGAAGGAAAAATACATAAAAATTATTTTGATGAAAGCCACAGCAGCT

TTATCAAATGCTTACGTTGCT**M**AATAGTAAAAAAAGCCACTTAAATTCCAAT
GGAAATTTTATACCCACATGTATTTATGTAAAACTTTTAAATAACATGTATTC
ATAATCACTTTTATATCCTCAACCAGTTTTTATGAAGCTAGAAAAAAATTCCT
TTATTaaagaaatgtaacattcaacaggt
Rev :5'-3': acctgttgaatgttacatttcttt

M272= EIF1A_Y STS4, (496 bp) **A to G** at position 212,
GROUP VIII
CaggaggggaccatgttttATAGTCCACAAAAACTCTGTTTAGATTATTCCTTCCTGGGA
CCCAGACCAATTTGTCTTCTTTTTACTTGCCTGTTGGCAGCATGGAATCTGTTT
CATTTTCTCTTTTTAGCTGTCACGACACACAGCTCTTGAGGTACTTGGTGACA
GTACAGTGCAGTCTTTCCTGGGCATTACTCTTTGCTCTCCCGAA**R**ACCCACTA
ACGGGTTGTGTGTATAATAAGGTTTTATTTTATTTTATTTTATTTTTTACTGCA
AAATTATTGGAGGATAAAGTGTATTCTGGGAGAAGTCTAATTAGAAAGAGTT
AGCAAAGGCTTATGCTTTTTCACTAACATTTTCTCAGATGGTACTGAACAACT
TCAGTAGGTATCTTGTTCTCACCTTTATTTCTAGTGATGAGATTCCCAGTTCTC
TAAGCCATCAGCTCTAAAGATCAGAGTATCTCCCTTTGCAaaatgtccattaaatctttgctg
For 5'-3'= caggaggggaccatgtttt
Rev 5'-3'=cagcaaagatttaatggacattt

M273= EIF1A STS8, (502 bp) **C to G** at position 189
GROUP II
CacatcaggaaaagggcatcCTTTGGCCTATACTTGTGAAGAGCTAGAGTAAGGTGCTC
CCCACCTTTGAGATTGCTAAAGTTGTCATTCTTTTGGAAATTTATGAGCTAAT
CATCATTTAGTCATTTGAAAAGCTGCCAAACTTTTGTAAAACCCAGTAAGGA
AAGCAGGTATGATCTTTGTCCTGA**S**GCAGCTAAGTTCAGGCACGATTAATTGC
TCGAAATATAGAATGTGTTTTCCTTTGTAGAAATTTAGTTTTGGCATGCCCTA
AAATGCATCAGAATCTGGATAAATCACAGAGTTCTGGAAGCCCAATTGTCTT
CTATAGTGGCACAGAACAATGTGAGACTGCCCCAGAGGTAGTGGGTGAATTC
AAGAAGTTAGATGTCTGGCTTTATGGTGGCCAGGTATATGTTTTATTCTATTT
GCAGTGTTAACATTTTTATTCAAATTCTTCAATCGATCCCTTAATATTACTGTA
atttgtagcctttctccctcc
For 5'-3'=cacatcaggaaaagggcatc
Rev 5'-3'=ggagggagaaaggctacaaat

M274= EIF1A_Y STS2a, (457 bp) **C to T** at position 47,
GROUPVIII w/M11
gccatgcccaagaataaagGTACTGCTGTAAGCCTCTGGGACTATA**Y**CTCGGCTTGCTCT
GCCAGTAACCCCGACGCCTGTTCCAGGCCGCAGTGACTGTTCTAACGGCGGT
ACTGGCCACTGCGACCCCAGCACTGTGTTCGGGAAAGGAGCTGGGAATGCCC
TATTTGGTCACATTGGGGTGGGACAGACGCCATTTTTGTGGGGCCTCCTTCGG
AAGATAGCGGGCTTTTGCTGCTGATTTCACGCCAGACGGAAAACGTATAGGT
AGGGACGGTTGAGGGACCTTAACCGGACGGCCTGGCTTTCCAGAATAGGCAC
ATG**S**AAACACTTCCCTGCTACTTTCCTGGAAGCGGTTCTTAACTTTGAAGACT
TACCTATCTGGACAGTTAAAAGTATTGCTAAGGATACTCCCTTTTCCTTGTTA
AACAGTGGGgaagccttgaagcatgtttag

For 5'-3'=gccatgcccaagaataaag
Rev 5'-3'=ctaaacatgcttcaaggcttc

M275= EIF1A_Y STS2b, (457 bp) **C to G** at position 325
GROUP X
gccatgcccaagaataaagGTACTGCTGTAAGCCTCTGGGACTATA**Y**CTCGGCTTGCTCT
GCCAGTAACCCCGACGCCTGTTCCAGGCCGCAGTGACTGTTCTAACGGCGGT
ACTGGCCACTGCGACCCCAGCACTGTGTTCGGGAAAGGAGCTGGGAATGCCC
TATTTGGTCACATTGGGGTGGGACAGACGCCATTTTTGTGGGGCCTCCTTCGG
AAGATAGCGGGCTTTTGCTGCTGATTTCACGCCAGACGGAAAACGTATAGGT
AGGGACGGTTGAGGGACCTTAACCGGACGGCCTGGCTTTCCAGAATAGGCAC
ATG**S**AAACACTTCCCTGCTACTTTCCTGGAAGCGGTTCTTAACTTTGAAGACT
TACCTATCTGGACAGTTAAAAGTATTGCTAAGGATACTCCCTTTTCCTTGTTA
AACAGTGGGgaagccttgaagcatgtttag
For 5'-3'=gccatgcccaagaataaag
Rev 5'-3'=ctaaacatgcttcaaggcttc

M276 EIF1A_Y STS12 (site b) (287 bp) **T to A** at position 58.
Group I associated mutation. Has another Group I site (M277) and a Group VI site (M267).
ttatcctgagccgttgtccctgTGTTTCCATTTCTCTTTTCCTCATTTCTCATCATC**W**ACATT
TCTCCTGTACTTGTTCATTAAATAATGATTCCTTGGATATACCAAGTCTGGAT
AGCGGATTCGATGGAAGCATTTTTGTAAATA**T**AC**G**TTCAGTATTTTGTGTGGA
AGAACACAATCTAGCTGATGCCTGCAATCCCAGCCCTTTGGAAAGCGAGGTG
GGTGGATTGCTTGAAGCTACGAGTTTGACACTAGCCTGGGCAACagggtacaaccgt
gtctctaca
newFor 5'-3'=ttatcctgagccgttgtccctg
Rev 5'-3=tgtagagacacggttgtaccct

M277 EIF1A_Y STS12 (site c) (287 bp) G to T at position.
Group I associated mutation. **G to T** at position 151 . Has another Group I site (M277) and a Group VI site (M267).
ttatcctgagccgttgtccctgTGTTTCCATTTCTCTTTTCCTCATTTCTCATCATC**T**ACATTT
CTCCT**G**TACTTGTTCATTAAATAATGATTCCTTGG**A**TATACCAAGTCTGGATA
GCGGATTCGAT**G**GAAGCATTTTTGTAAATA**T**AC**K**TTCAGTATTTTGTGTGGA
AGAACACAATCTAGCTGATGCCTGCAATCCCAGCCCTTTGGAAAGCGAGGTG
GGTGGATTGCTTGAAGCTACGAGTTTGACACTAGCCTGGGCAACagggtacaaccgt
gtctctaca
newFor 5'-3=ttatcctgagccgttgtccctg
Rev 5'-3'=tgtagagacacggttgtaccct

M278= DBY int12n, site c ((nominal, 418 bp)) **T to G** at position 374, Site within STS with 7 T homopolymer.
Group I.
aaatattgcatctggctggaATTGCATCAAAGGTTTATTAACTGCCTTAAGGAGAGTTGGC
AATATTTTAGTATTTGAGGGGATGGAAGAGA**C**CTTAAACATCTAACTTCCTA

AATCTGGGAAGTACAATCGATTTAGTACAATAGATCTAGATTTAGGAAGTAC
AATTATTCATTTGTCTAATATTGGAGATTTAAAAGCAGGGGAAAATAACTTTA
TTAACTTGTAACTTTAAACATTCATTGAAATGTTTGAATTTAGGTAAGTGTGT
GGTTGTGG**G**Agtgagtttactcttgtca**ttTTTTTTTT**ATCAGTTTGTAGACATGGAAAGTA
GGCAACAATGAGGG**TTTTTTT**GTTTTAACACAAGTATACCT**K**ATTCTTAACG
AGCATATTaagattacatagttacttttggactt
For 5'-3'=aaatattgcatctggctgga
Rev 5'-3'=aagtccaaaagtaactatgtaatctt
New Rev 5'-3'=aatgacaagagtaaactcac to exclude poly T region

M279= DBY int12n, site d ((nominal, 418 bp)) **C to T** at position 93, Site within STS with 7 T homopolymer.
Group I
aaatattgcatctggctggaATTGCATCAAAGGTTTATTAACTGCCTTAAGGAGAGTTGGC
AATATTTTAGTATTTGAGGGGATGGAAGAGA**Y**CTTAAACATCTAACTTCCTA
AATCTGGGAAGTACAATCGATTTAGTACAATAGATCTAGATTTAGGAAGTAC
AATTATTCATTTGTCTAATATTGGAGATTTAAAAGCAGGGGAAAATAACTTTA
TTAACTTGTAACTTTAAACATTCATTGAAATGTTTGAATTTAGGTAAGTGTGT
GGTTGTGG**G**Agtgagtttactcttgtca**ttTTTTTTTT**ATCAGTTTGTAGACATGGAAAGTA
GGCAACAATGAGGG**TTTTTTT**GTTTTAACACAAGTATACCT**T**ATTCTTAACG
AGCATATTaagattacatagttacttttggactt
For 5'-3'=aaatattgcatctggctgga
Rev 5'-3'=aagtccaaaagtaactatgtaatctt
New Rev 5'-3'=aatgacaagagtaaactcac to exclude poly T region

M280 revised B9.36 c (386 bp) STS **G to A** at position 280
Group VI
ccagtcagcagtacaaaagttgACAGCTTCAGCAAAATTGTAGCCTTGGTTAAAACCACTG
TGGTAAGCACGAGGAAAAGTGATGACAAACTCCCCTGCACACTGGTTTGTGC
GGACAACCTAAAAAGGAGAAAAAAGCAGAAAGAGGTGTGGGTCAGAACTAA
TGGGCCAGATGTGAACTCAAAGATGTCTCTAGATGCTGTAACAGATGTAGGA
AG**A**GTGGAAAGGCTCTATCTTCAAGTACGTGTCCTAAAAGAAAATGAGATTG
TGAATTTAAAA**R**TGGTATTCATAGAAAAGTACTCAAAATATGTGTAATTCAA
AAAACAA**A**TATAGAGGGGTCCACGAACAAGTGAAAAGACTCTttgcttctataatcaaa
gaaatgc
newFor 5'-3' = ccagtcagcagtacaaaagttg
newRev 5'-3' = gcatttctttgattatagaagcaa

M281 = G3.27f (393 bp) **G to A** at position 247.
Discovered while typing M123
tggtaaactctacttagttgcctttTGGAAATGAATAAATCAAGGTAGAAAA**G**CAATTGAGAT
ACTAATTCA**T**GCTCTCAGGGGAAAATCTGAATAAAGCTATCTTTTCTAACACA
GAGCAAGTGACTCTCAAAGTCACAGTATCTGAACTAGCATATCA**G**CATCGCC
TGAATACCTAGAAATGCAAATTCCTGGGCAACACCAGAATCTAACAAAGCAA
AAAACTATGGGGGGAACAGGGAAGT**C**R**G**TTTAATAATACTGAGTTTGTGCA
ACCTCAACTTTGCTTTA**T**AGGAAAGCAAAATCTCAATATGATAAAGTTTTCTT

CAACAAAACTCTGAGATAACTATGTTGAGGGAAAGAAGTTGATCACATgcaaga
aaatctaattcgctg
For = tggtaaactctacttagttgccttt
Rev 5'-3' = cagcgaattagattttcttgc

M282 = G3.27g (393 bp) **A to G** at position 316.
Group VI
tggtaaactctacttagttgcctttTGGAAATGAATAAATCAAGGTAGAAAA**G**CAATTGAGAT
ACTAATTCA**T**GCTCTCAGGGGAAAATCTGAATAAAGCTATCTTTTCTAACACA
GAGCAAGTGACTCTCAAAGTCACAGTATCTGAACTAGCATATCA**G**CATCGCC
TGAATACCTAGAAATGCAAATTCCTGGGCAACACCAGAATCTAACAAAGCAA
AAAACTATGGGGGGAACAGGGAAGT**CG**GTTTAATAATACTGAGTTTGTGCAA
CCTCAACTTTGCTTTA**T**AGGAAAGCAAAATCTCAATATGATAA**R**GTTTTCTTC
AACAAAACTCTGAGATAACTATGTTGAGGGAAAGAAGTTGATCACATgcaagaa
aatctaattcgctg
For = tggtaaactctacttagttgccttt
Rev 5'-3' = cagcgaattagattttcttgc

M283 = DBY STS 09b (429 bp) **A to G** at position ?
STS also contains M200.
ggcttacacttgcagactttgCAAATCTTAAGACTAACAAATCCTTGAAATCACACAGCTT
GCAAATACGTACTAAACTGCACAAGGTGTGTGTTCTATATGTGCAGTTTTAGC
GTATTTTAGTTGCATAGGTTTCCATGGTATTTATAGTCTCTTGTGCTAAATTTG
GCCAAAGATGATTGTCCACCACTAAAAATGCCTCTCCCACTTGGAATTCTGTA
CTGATTTTGTGGCCAGATGCAATGATCTTTAAAAACAAATCTTTTCAATGGCA
TAAGAAGTTGAC**R**AAAATTTCTTAAAGTGCAATAGATTTTCAA**G**TGTATTGTG
CCTTGTTCTAAAACTTTTAAGTAGGTGCACTTGACAGTATTGAGGTCATTTGT
TAAGGTGCTATTTCAATTAGTGTAggtttagactcttgtacatttctcc
For = ggcttacacttgcagactttg
Rev: 5'-3'= ggagaaatgtacaagagtctaaacc

M284 = EIF1AY STS34a, (399 bp nominal) **del ACAA** at position 105, STS has another marker, M306,
GroupIX.
GgcagttttcatttaagcagaGGCAACAAATGTAATACTAATGTTTGATTATTATAGAAAA
AAGTATTCATCTTAGCAAAGTTTTAACTATGGGATTATTTTTA**ACAA**ACAAT
TGTGTTTTCTTTTTCTTAAAGACAAACACAATGCATACTTACTGCCGAAAGCT
TGACAAGATTAAAATAAGTCCCTCATGACACCATCAAAGAGAATATGCACTG
TTGTAAAGCCTG**C**GTATTTTACTTGGCAGCTATTTTCATTATTTATCATATTGC
ATTTTATGAAAAGATTTTTATATAAACATGAAGATCTTGATGAAATTATTGGC
ATTTCAGGAAGTGCTGAAATGTTATTGGAAGTGATGAAATTATTGGCATTTCA
Ggaagtgctgaaagtttcgct
F 5'-3' = ggcagttttcatttaagcaga
R 5'-3' = agcgaaactttcagcacttc

M285 EIF1A_Y STS12 (site d) (287 bp) **G to C** at position 70

(GroupVI)
ttatcctgagccgttgtccctgTGTTTCCATTTCTCTTTTCCTCATTTCTCATCATC**T**ACATTT
CTCCTGTACTTGTTCATTAAATAATGATTCCTTGG**A**TATACCAAGTCTGGATA
GCGGATTCGATGGAAGCATTTTTGTAAATA**T**AC**G**TTCAGTATTTTGTGTGGAA
GAACACAATCTAGCTGATGCCTGCAATCCCAGCCCTTTGGAAAGCGAGGTGG
GTGGATTGCTTGAAGCTACGAGTTTGACACTAGCCTGGGCAACagggtacaaccgtgt
ctctaca
newFor 5'-3'=ttatcctgagccgttgtccctg
Rev 5'-3'=tgtagagacacggttgtaccct

M286 EIF1A_Y STS12 (site e) (287 bp) **G to A** at position 129.
(GroupVI)
ttatcctgagccgttgtccctgTGTTTCCATTTCTCTTTTCCTCATTTCTCATCATC**T**ACATTT
CTCCT**G**TACTTGTTCATTAAATAATGATTCCTTGG**A**TATACCAAGTCTGGATA
GCGGATTCGAT**R**GAAGCATTTTTGTAAATA**T**AC**G**TTCAGTATTTTGTGTGGA
AGAACACAATCTAGCTGATGCCTGCAATCCCAGCCCTTTGGAAAGCGAGGTG
GGTGGATTGCTTGAAGCTACGAGTTTGACACTAGCCTGGGCAACagggtacaaccgt
gtctctaca
newFor 5'-3'=ttatcctgagccgttgtccctg
Rev 5'-3'=tgtagagacacggttgtaccct

M287 EIF1A_Y STS12 (site f) (287 bp) **A to T** at position 100. This is one of 3 M201 related mutations.
(Group VI)
ttatcctgagccgttgtccctgTGTTTCCATTTCTCTTTTCCTCATTTCTCATCATC**T**ACATTT
CTCCT**G**TACTTGTTCATTAAATAATGATTCCTTGG**W**TATACCAAGTCTGGAT
AGCGGATTCGAT**G**GAAGCATTTTTGTAAATA**T**AC**G**TTCAGTATTTTGTGTGGA
AGAACACAATCTAGCTGATGCCTGCAATCCCAGCCCTTTGGAAAGCGAGGTG
GGTGGATTGCTTGAAGCTACGAGTTTGACACTAGCCTGGGCAACagggtacaaccgt
gtctctaca
newFor 5'-3'=ttatcctgagccgttgtccctg
Rev 5'-3'=tgtagagacacggttgtaccct

M289 = B9.36new d (386 bp) **G to A** at position 227 Group VI.
ccagtcagcagtacaaaagttgACAGCTTCAGCAAAATTGTAGCCTTGGTTAAAACCACTG
TGGTAAGCACGAGGAAAAGTGATGACAAACTCCCCTGCACACTGGTTTGTGC
GGACAACCTAAAAAGGAGAAAAAAGCAGAAAGAGGTGTGGGTCAGAACTAA
TGGGCCAGATGTGAACTCAAAGATGTCTCTAGATGCTGTAACAGATGTAGGA
AG**A**GTGGAAA**R**GCTCTATCTTCAAGTACGTGTCCTAAAAGAAAATGAGATTG
TGAATTTAAAA**G**TGGTATTCATAGAAAAGTACTCAAAATATGTGTAATTCAA
AAAACA**A**ATATAGAGGGGTCCA**G**GAACAAGTGAAAAGACTCTttgcttctataatcaaa
gaaatgc
For 5'-3' = ccagtcagcagtacaaaagttg
Rev 5'-3' = gcatttctttgattatagaagcaa

M290 = B9.36new e (386 bp) **G to A** at position 343. Group III

ccagtcagcagtacaaaagttgACAGCTTCAGCAAAATTGTAGCCTTGGTTAAAACCACTG
TGGTAAGCACGAGGAAAAGTGATGACAAACTCCCCTGCACACTGGTTTGTGC
GGACAACCTAAAAAGGAGAAAAAAGCAGAAAGAGGTGTGGGTCAGAACTAA
TGGGCCAGATGTGAACTCAAAGATGTCTCTAGATGCTGTAACAGATGTAGGA
AG**A**GTGGAAA**G**GCTCTATCTTCAAGTACGTGTCCTAAAAGAAAATGAGATTG
TGAATTTAAAA**G**TGGTATTCATAGAAAAGTACTCAAAATATGTGTAATTCAA
AAAACA**A**ATATAGAGGGGTCCA**Y**GAACAAGTGAAAAGACTCTttgcttctataatcaaa
gaaatgc
newFor 5'-3' = ccagtcagcagtacaaaagttg
newRev 5'-3' = gcatttctttgattatagaagcaa

M291 = EIF1AY STS16, (480 bp) **A to G**, at position 358,
(Group III)
cggagtctggctttgttggcCAGGTTGGAGTGCAGTGGCATGATCTCGGCTCAGGGCAAT
GTCCGTCTCCTGGACTCAAGCAGTTCTCCTGCCTCAGCCTCCCCAGTAGCTGG
GATTAGAGGTGTGTGACACCATGCCCGGCTAATTTTTGTATTTTTAGTAGAGA
TGGGGTTTCACCATGTTGGCCAGGCTGGTCTCGAACTCCTGACCTCAGGTAAT
GCACCCGCCTCGGCCTCCCAAAGTGGTGGGATTATAGGCGTGAGTAACCATG
CCTGGCCTTTCACTCTTATTTTCTAAGAACTTTAGAATAATCACCGAGATATT
CTAAAGTAAACAGGAATTTTTAATGGTTAAGCT**R**TTATTTGTCTTTGTCATTTC
TGAGTTTAGGGATAGTGAAGATAGAGTTAGGCCTCATGTGTGAGAGACTGAT
GTAGCATTATAGTGTATATTTTGAAATGTGccaccgtgatgttcaaaagt
For = cggagtctggctttgttggc
Rev 5'-3' = acttttgaacatcacggtgg

M292 = EIF1AY STS19, (556 bp) **A to G**, at position 373.
Group III
TttaacaaatgtggaccaagaTCTCAACCTTTTTTTTTATctcctctcctcagagtatgcTCAGGTAAT
CAAAATGTTGGGAAATGGACGATTGGAAGCATTGTGTTTTGATGGTGTAAAG
AGGTTATGCCATATCAGAGGGAAATTGAGAAAAAAGGTAGGTGTGTAGGTTA
CTTTTCAATAAAAATTTGCCGCAAAAAATGTCTCTGCTTTAAATACATGGTCC
AAGCAATTTATTTTTGTGAGTTCCCAAAATAATTTATACAGCAATGATTCATG
TGACAATGTGAATAAATAGAAAAAGTCTTTGATAACTTTTAGATTTACTTTTA
AAGAATAATTTGTTTGTTTAACTTCTGTTGTATTCCTACC**R**GAAATGTTTACTC
TGATATTAGTATTGAAGAAACCAGACAAATCTAATATATAACACAAATGGTC
TTGACTCAGATGTTAATGCTGTGAAAGAATGAAAAATCTGGGAATTACTTTA
GCTTAAAAGAGATTGATCGGTGCATATCCCTTTGTTAGGTTTTGgattgggggaaata
gttttagg
Original F 5'-3' = tttaacaaatgtggaccaaga
Rev 5'-3' = acttttgaacatcacggtgg

M293 = EIF1AY STS20a, (507bp) **T to G**, at position 299.
Group III. STS also contains **M294**
CatggtccaagcaatttatttttgTGAGTTCCCAAAATAATTTATACAGCAATGATTCATGTG
ACAATGTGAATAAATAGAAAAAGTCTTTGATAACTTTTAGATTTACTTTTAAA
GAATAATTTGTTTGTTTAACTTCTGTTGTATTCCTACCAGAAATGTTTACTCTG

ATATTAGTATTGAAGAAACCAGACAAATCTAATATATAACACAAATGGTCTT
GACTCAGATGTTAATGCTGTGAAAGAATGAAAAATCTGGGAATTACTTTAGC
TTAAAAGAGATTGATCGGTGCATA**K**CCCTTT**T**GTTAGGTTTTGGATTGGGGGA
AATAGTTTTAGGTGGTACTAGGAAAATTGGAATATGGAATATGTTAGAAACT
CTATTTGTTAGTAATACCACATCAGGTAGTTT**T**ATAAATTACACTGATTAAAA
GTCTCTACTACTCAGATTTTTAATTAAAATAATAAAAACTTATTTTTGG**C**TGA
Gctctgtggaagtattagccagc
F 5'-3' = catggtccaagcaatttatttttg
R 5'-3' = gctggctaatacttccacagag

M294 = EIF1AY STS20b, (507bp) **C to T**, at position 305
Catggtccaagcaatttatttttg**T**GAGTTCCCAAAATAATTTATACAGCAATGATTCATGTG
ACAATGTGAATAAATAGAAAAAGTCTTTGATAACTTTTAGATTTACTTTTAAA
GAATAATTTGTTTGTTTAACTTCTGTTGTATTCCTACCAGAAATGTTTACTCTG
ATATTAGTATTGAAGAAACCAGACAAATCTAATATATAACACAAATGGTCTT
GACTCAGATGTTAATGCTGTGAAAGAATGAAAAATCTGGGAATTACTTTAGC
TTAAAAGAGATTGATCGGTGCATA**T**CCCTT**Y**GTTAGGTTTTGGATTGGGGGA
AATAGTTTTAGGTGGTACTAGGAAAATTGGAATATGGAATATGTTAGAAACT
CTATTTGTTAGTAATACCACATCAGGTAGTTT**T**ATAAATTACACTGATTAAAA
GTCTCTACTACTCAGATTTTTAATTAAAATAATAAAAACTTATTTTTGG**C**TGA
Gctctgtggaagtattagccagc
F 5'-3' = catggtccaagcaatttatttttg
R 5'-3' = gctggctaatacttccacagag

M295 = EIF1AY STS20c, (507bp) **T to C**, at position 411,
(Group VIII). STS also contains M294 mutation
catggtccaagcaatttatttttg**T**GAGTTCCCAAAATAATTTATACAGCAATGATTCATGTG
ACAATGTGAATAAATAGAAAAAGTCTTTGATAACTTTTAGATTTACTTTTAAA
GAATAATTTGTTTGTTTAACTTCTGTTGTATTCCTACCAGAAATGTTTACTCTG
ATATTAGTATTGAAGAAACCAGACAAATCTAATATATAACACAAATGGTCTT
GACTCAGATGTTAATGCTGTGAAAGAATGAAAAATCTGGGAATTACTTTAGC
TTAAAAGAGATTGATCGGTGCATA**T**CCCTT**T**GTTAGGTTTTGGATTGGGGGAA
ATAGTTTTAGGTGGTACTAGGAAAATTGGAATATGGAATATGTTAGAAACTC
TATTTGTTAGTAATACCACATCAGGTAGTTT**Y**ATAAATTACACTGATTAAAAG
TCTCTACTACTCAGATTTTTAATTAAAATAATAAAAACTTATTTTTGG**C**TGAGc
tctgtggaagtattagccagc
F 5'-3' = catggtccaagcaatttatttttg
R 5'-3' = gctggctaatacttccacagag

M296 = EIF1AY STS21=STS20d, (536 bp) **C to T**, at position 165,
(Group VIII)
gattgggggaaatagttttaggTGGTACTAGGAAAATTGGAATATGGAATATGTTAGAAAC
TCTATTTGTTAGTAATACCACATCAGGTAGTTTTATAAATTACACTGATTAAA
AGTCTCTACTACTCAGATTTTTAATTAAAATAATAAAAACTTATTTTTGG**Y**TG
AGCTCTGTGGAAGTATTAGCCAGCATACACCTGTAGTCCCAGCTACTGAGGA
GGCTGAGCCCAGGAGTTCAAGGTTCCCATGAGCTAAAAATTGTGCTAATGCT

CTCCAGTCTGGGTGATAGAGCGAATCTCTATCTCAAAAAGAAAAAAAAAAAA
ATCTTTCTGGTATGTTAACATTCTTTCTTTTCCAAATTAGTGGCATTTTAGGGA
TTCTCTTAGTCCATTTGGGCTGTCACTGACTGGGTAGATTATAAAAAGCAGAA
ATTTTATTTCTCATAGTTTTGGAGAAAGAGAAATCTATTTAATATTTGGTGAG
GACCCATTTCCTGATTATTATGTGGTGCCTTctggcttagtccacacatagtg
F 5'-3' = gattgggggaaatagttttagg
R 5'-3' = cactatgtgtggactaagccag

M297 = EIF1AY STS24, (506 bp) **A to G**, at position 303,
(Group VII)
TtggttggtctacgggactATCAGGTAAAAATAACATTTAAAGTTGTGGTATGTCTGTGT
TTAAGCAGTTGTTAATGTTTGGAAGGTAACTATACTAGCATCTTTGACCCATT
CCAGCCCAGGTTGCTTTCTCACCATTCTGCCTGCCATCATCATTTATTAAGGG
CCAGTTGTATTTCAGACTATAGTATTTTTCAAATTTGACATAATTCTCACTGAT
AGTAAATGGTACATATATTTTTGTGGAAAGACATAAAGTTTTTAATTCTTTGT
TTTCATTGTTAATATAATGTGCAGTAAAT**R**TTTTCTTGCAGGCTTGGGCAAGT
ACTGTAGACCATCTGTCCTCATCCATTTAAAGGCCAATGGTGTTTCAGGCATT
CAGCTAGGTATTTCAGACATTGTAGTTCCCAAATGCCGGTCTGTTAAATAGTA
TTGGTGCAGGCTGAATTTTCAGTGCTCTGAAGTCAAATTAGAAGATACATAGT
Tacgatgtttttcatggagca
F 5'-3' = ttggttggtctacgggact
R 5'-3' = tgctccatgaaaaacatcgt

M298 = EIFIA STS 27 (445 bp) **G to A** at position 230,
Group II
AaataccattttcataatttccttAATATTTTTAGACATTATTTCTTTTTAAGTCTTAGATAAA
CTAAGTCCAACTTCTGGGATTCCTCAGGAATAGTATTTTTTTTTTCCCTGTGTT
TGAGCCACTTTTTTAAATCTTTTTTTTTTTTTTAAACCGAACAATTTAACTACA
ACATAGCAGTTCTGGAAATCAGATTGCTGCCTCTCGGGGCTGTTGTTGATACT
GCTT**R**TTTGGTGACTTTTCTGAACTAATTCTTTGGCCATTGAATAGTTGGTTA
GTTTAGTGGGCAGTTCATGTTTGAACTAAGATTTCATTAAAACCACCAAGAAT
TTAATCATTTAAAGAGGAATCTTGTACATGTAGAGGAATACTTTGAGCATTCA
GCCAATGTTGGTAAACTGACACCTCTTCCTTAGTCTTCATTtcttgctgtgcaggatctca
Original F 5'-3' = aaataccattttcataatttcctt
Original R 5'-3' = tgagatcctgcacagcaaga

M299 = EIF1AY STS29, (483 bp) **T to G**, at position 127,
Group I
CggacttggtctgtgcttttcAGTAGCTGCTATTGTGTTGGTTTTTATTAAACTGAGGTAAG
GAATGGGAATAGGGGAACTTAAAAGCCCACACTGCTTTTTCTTAGTAAGGTT
CACCTATTTTTC**K**TGAATAAACGCTCCTTAGTGTTTATTGCATTCATTTGGTTA
ATTTTCAGATTTCTGATATATGGATTTTGACCATGTTTGTCAATGTTCTTATTT
CTTTTCTGAAGGAACAAATTTTAGCAAGTCCTTATTCTGCCATTCCTGCAATC
ACTGCAAGAAAGCATTTATTTTGATAAGACTTAATTACACATTGACTTTGTTT
CTTTTTCATATATCAAATAAAAAGTTGTACTGTGCTTTTAAAATGTTATTTTTA

TGTCCATTATATTATTCGAATTATCATTTTAACAAAAACTGGTTTGCACATTA
CAGTTTGAAAAGTGTTGGTCTATTTCATactgccattgtgacagatca
F 5'-3' = cggacttggtctgtgcttttc
R 5'-3' = tgatctgtcacaatggcagt

M300 = EIF1AY STS31, (500 bp) **G to A** at position 153,
STS also contains **M301**, Group III
CaggcaggtctactttcaatctTAAGGAAGTAGGTATGTATTTTTAAAATCAAGCTATTTTT
CAAGTTCCATAGACAATTCTGTTAGATAATCTATACTAAGAACTACTGATGCA
TAGAAAAGTTTATTATTGTTGTTTTTGTTTTTTTGAA**R**GAGTTTCGCTCTGTTG
CCCAGGCTGGAGTGCAGTGGCTTGATCTCGGCTCACTGCAAGCTGCGCCTCCT
GGGTTCATGCCATTCTCCTGCCTCAGCCTCCTGAGTAGCTGGGACTACAGATG
CCTGCCACCACGCCCAGCTAATTTTTTGTATTTTTAGTAGAGATGGGGTTTCA
TCATGTTAGCC**A**GTATGGTCTCGATCTCCTGACCTCATGATCCGCCCGCCTTG
GCCTCCCAAAGTGCTGGGATTACAGGCGCGAGCCACCGTGCCTGGCCTAGAA
AAGTGTATTACCTTTTTAACATCATTATTCTTTACTCCATTTTTAgttttgaattgcagtgt
ttgac
F 5'-3' = caggcaggtctactttcaatct
R 5'-3' = gtcaaacactgcaattcaaaac

M301 = EIFIA STS 31 (500 bp) **A to C** at position 340bp.
(Group III) STS also contains **M300**, a GroupVII marker
CaggcaggtctactttcaatctTAAGGAAGTAGGTATGTATTTTTAAAATCAAGCTATTTTT
CAAGTTCCATAGACAATTCTGTTAGATAATCTATACTAAGAACTACTGATGCA
TAGAAAAGTTTATTATTGTTGTTTTTGTTTTTTTGAA**G**GAGTTTCGCTCTGTTG
CCCAGGCTGGAGTGCAGTGGCTTGATCTCGGCTCACTGCAAGCTGCGCCTCCT
GGGTTCATGCCATTCTCCTGCCTCAGCCTCCTGAGTAGCTGGGACTACAGATG
CCTGCCACCACGCCCAGCTAATTTTTTGTATTTTTAGTAGAGATGGGGTTTCA
TCATGTTAGCC**M**GTATGGTCTCGATCTCCTGACCTCATGATCCGCCCGCCTTG
GCCTCCCAAAGTGCTGGGATTACAGGCGCGAGCCACCGTGCCTGGCCTAGAA
AAGTGTATTACCTTTTTAACATCATTATTCTTTACTCCATTTTTAgttttgaattgcagtgt
ttgac
F 5'-3' = caggcaggtctactttcaatct
R 5'-3' = gtcaaacactgcaattcaaaac

M302 = EIFIA STS 32a (527bp) **A to G** at position 230
(Group VII)
CaaagtgctgggattacaggCGCGAGCCACCGTGCCTGGCCTAGAAAAGTGTATTACCT
TTTTAACATCATTATTCTTTACTCCATTTTTAGTTTTGAATTGCAGTGTTTGAC
CTTAAAAGTTTTATATTACAATTTTTTTAATTAGTCTTTTATTTTTTCCAAGAG
ACTTCTAATTAAAAGGGAATAGTAAATAAAAGCACTGTGCTTGCCTTTTGTGC
TTTTATTAA**R**GTGAAATCTCTACAATCTTTCCTAAGCTGTTAATCACTGTTTA
CTAATGAACATAAACCACTTCCTAATTATTCAGACTCAAGAATTTTTTTCTAG
AGGGTATTGGGGTAGGCAAAGAAAA**G**CAGGAGAGTTTGTAACAAACAGTAT
GTGGGATTTTTTTAGATGTGTTCAATTTGAAAGTAACTTGTGA**A**ACAACTGGT

GATATTTTGGTATAAGACGTTTTGAAAGTTATTTGTTTATTTCTAAGGATAAC
AAAGCTGATGTAATTTTAAAGTacaatgcagatgaagctagaag
F 5'-3' = caaagtgctgggattacagg
R 5'-3' = cttctagcttcatctgcattgt

M303 = EIFIA STS 32b (527bp) **G to C** at position 352,
(Group X)
CaaagtgctgggattacaggCGCGAGCCACCGTGCCTGGCCTAGAAAAGTGTATTACCT
TTTTAACATCATTATTCTTTACTCCATTTTTAGTTTTGAATTGCAGTGTTTGAC
CTTAAAAGTTTTATATTACAATTTTTTTAATTAGTCTTTTATTTTTTCCAAGAG
ACTTCTAATTAAAAGGGAATAGTAAATAAAAGCACTGTGCTTGCCTTTTGTGC
TTTTATTAAA**G**TGAAATCTCTACAATCTTTCCTAAGCTGTTAATCACTGTTTAC
TAATGAACATAAACCACTTCCTAATTATTCAGACTCAAGAATTTTTTTCTAGA
GGGTATTGGGGTAGGCAAAGAAAA**S**CAGGAGAGTTTGTAACAAACAGTATG
TGGGATTTTTTTAGATGTGTTCAATTTGAAAGTAACTTGTGA**A**ACAACTGGTG
ATATTTTGGTATAAGACGTTTTGAAAGTTATTTGTTTATTTCTAAGGATAACA
AAGCTGATGTAATTTTAAAGTacaatgcagatgaagctagaag
F 5'-3' = caaagtgctgggattacagg
R 5'-3' = cttctagcttcatctgcattgt

M304 = EIFIA STS 32c (527bp) **A to C** at position 421
CaaagtgctgggattacaggCGCGAGCCACCGTGCCTGGCCTAGAAAAGTGTATTACCT
TTTTAACATCATTATTCTTTACTCCATTTTTAGTTTTGAATTGCAGTGTTTGAC
CTTAAAAGTTTTATATTACAATTTTTTTAATTAGTCTTTTATTTTTTCCAAGAG
ACTTCTAATTAAAAGGGAATAGTAAATAAAAGCACTGTGCTTGCCTTTTGTGC
TTTTATTAAA**G**TGAAATCTCTACAATCTTTCCTAAGCTGTTAATCACTGTTTAC
TAATGAACATAAACCACTTCCTAATTATTCAGACTCAAGAATTTTTTTCTAGA
GGGTATTGGGGTAGGCAAAGAAAA**G**CAGGAGAGTTTGTAACAAACAGTATG
TGGGATTTTTTTAGATGTGTTCAATTTGAAAGTAACTTGTGA**M**ACAACTGGT
GATATTTTGGTATAAGACGTTTTGAAAGTTATTTGTTTATTTCTAAGGATAAC
AAAGCTGATGTAATTTTAAAGTacaatgcagatgaagctagaag
F 5'-3' = caaagtgctgggattacagg
R 5'-3' = cttctagcttcatctgcattgt

M305 = EIFIA STS 33 (545 bp) **C to T** at position 331
(Group I)
AacttgtgaaacaactggtgatATTTTGGTATAAGACGTTTTGAAAGTTATTTGTTTATTTC
TAAGGATAACAAAGCTGATGTAATTTTAAAGTACAATGCAGATGAAGCTAGA
AGCCTGAAGGCATATGGCGAGCTTCCAGAACATGGTAAGATCAAAATGATTT
TATCTCCTCATTATTTGATATTAATGTTTGTTGGTATTTAGGTGAAGGTATTTC
CGTAGAACTCTTGTTTTACATACTGTTTTAGTGTATACTTAAAAATTTGTTATA
AGTAGTCTTGCCTATACTTCAGTTTACTTATGATACTTTGGAAAAGATATTAA
TAA**Y**TGGAAATCTCTAATAAAAACGTTATGAACTTGAAAGTAGAAGTCTCTA
ATAAAGAGATTATGAATTATGAAAGTTCCTTTAGTGACAACTTTATAAATTCA
TAAGCTCTGGATTTGTATATAAGATCTGTCAAAGAAATACGTTTTTTATAGTG
TTTTTCTAAACAGTTCTCAAGACTGGCAGTTTTCATTTaagcagaggcaacaaatgtaat

F 5'-3' = aacttgtgaaacaactggtgat
R 5'-3' = attacatttgttgcctctgctt

M306 = EIFIA STS 34b (399 bp) **C to A** at position 231.
Group IX. STS also contains **M284**, a Group VI marker.
GgcagttttcatttaagcagaGGCAACAAATGTAATACTAATGTTTGATTATTATAGAAAA
AAGTATTCATCTTAGCAAAGTTTTAACTATGGGATTATTTTTA**A**C**AA**ACAATT
GTGTTTTCTTTTTCTTAAAGACAAACACAATGCATACTTACTGCCGAAAGCTT
GACAAGATTAAAATAAGTCCCTCATGACACCATCAAAGAGAATATGCACTGT
TGTAAAGCCTG**C**GTATTTTACTTGGCAGCTATTTTCATTATTTATCATATTGC
ATTTTATGAAAAGATTTTTATATAAACATGAAGATCTTGATGAAATTATTGGC
ATTTCAGGAAGTGCTGAAATGTTATTGGAAGTGATGAAATTATTGGCATTTCA
Ggaagtgctgaaagtttcgct
F 5'-3' = ggcagttttcatttaagcaga
R 5'-3' = agcgaaactttcagcacttc

M307 = EIFIA STS 35 (500 bp) **G to A** at position 282
(Group VI)
TtattggcatttcaggaagtgCTGAAATGTTATTGGAAGTGATGAAATTATTGGCATTTCA
GGAAGTGCTGAAAGTTTCGCTTTCATTACTTGGGGATAAGCATGATCATGATT
TAACCAAGTATTTCTCACTGATTTGATAAGTCTGTTTAAATAATTGGTTAACT
AGTTGTTGTAATTTCAAGAGAACTTTATGTATTTTGAGGATAAGTTGTTAACC
TGTGCTCAAATCCTTTTTGAAGGCTACATGGAAATGGTTGGCTATTGAGTTAG
CATAATCA**R**TCTGCCTACCATACTTAAAGTACCTTTTGTATATGTGCTAAGTG
AGAATTAAAAATACCTTTTAAAAACAAATGAAAAATACAGCACAATACAGCA
CATTCGTTCTTTGTTTTTTGAAACAGAGTCTTGCTCTGTCACCCAGGCAGGAG
TGCAGTGGCACCATCTCAGCTCCCTGCATTCTACGCCTGCCAAGTTCAAgctatttt
cctgcctcaccc
F 5'-3' = ttattggcatttcaggaagtg
R 5'-3' = gggtgaggcaggaaaatagc

M308 = EIFIA STS 37a (444 bp) **T to C** at position 70
(Group I)
AaactttacagtcctttgggataGTATTTACTGCAAAAATCAATTTTAGCTTCGGCAGTAGG
CACTTCA**Y**AATCAACGTTAAGTAAGAGTGTCTAAAGAGATAGTTTTGAGAAC
ACGTCCTCTATTAAGAGAAATGCTTAGTATGTTAAAAGAAGAATTTTGTTTGA
ACCAGTTTGATGCAGCACTGAAATTACAACAT**A**CTTCAAAGGTTTGTTAAAAT
GAAGGGCCTGTTGCCAGGACATGTAATAGAATTACATGGTTGAGCATCAGTT
TGTACTGGCCAGACTCTTGTTTTGGAGTTAGTTTGTGCTTATTTTGTGGAAATG
ATTGTTTTTCCTAGTAACAAAGCAG**C**GCAGTTCACAAAGCAGTAAATGCTTC
AGCTCTCTTTTTCAGTTAACTATATTGAAATTAAATTCACTTTgatttttcttccctctcttg
aga
F 5'-3' = aaactttacagtcctttgggata
R 5'-3' = tctcaagagagggaagaaaaatc

M309 = EIFIA STS 37b (444 bp) **A to G** at position 200

(Group II)
AaactttacagtcctttgggataGTATTTACTGCAAAAATCAATTTTAGCTTCGGCAGTAGG
CACTTCA**T**AATCAACGTTAAGTAAGAGTGTCTAAAGAGATAGTTTTGAGAAC
ACGTCCTCTATTAAGAGAAATGCTTAGTATGTTAAAAGAAGAATTTTGTTTGA
ACCAGTTTGATGCAGCACTGAAATTACAACAT**R**CTTCAAAGGTTTGTTAAAA
TGAAGGGCCTGTTGCCAGGACATGTAATAGAATTACATGGTTGAGCATCAGT
TTGTACTGGCCAGACTCTTGTTTTGGAGTTAGTTTGTGCTTATTTTGTGGAAAT
GATTGTTTTTCCTAGTAACAAAGCAG**C**GCAGTTCACAAAGCAGTAAATGCTT
CAGCTCTCTTTTTCAGTTAACTATATTGAAATTAAATTCACTTTgatttttcttccctctctt
gaga
F 5'-3' = aaactttacagtcctttgggata
R 5'-3' = tctcaagagagggaagaaaaatc

M310 = EIFIA STS 37c (444 bp) **C to T** at position 352
(Group III)
AaactttacagtcctttgggataGTATTTACTGCAAAAATCAATTTTAGCTTCGGCAGTAGG
CACTTCA**T**AATCAACGTTAAGTAAGAGTGTCTAAAGAGATAGTTTTGAGAAC
ACGTCCTCTATTAAGAGAAATGCTTAGTATGTTAAAAGAAGAATTTTGTTTGA
ACCAGTTTGATGCAGCACTGAAATTACAACATACTTCAAAGGTTTGTTAAAAT
GAAGGGCCTGTTGCCAGGACATGTAATAGAATTACATGGTTGAGCATCAGTT
TGTACTGGCCAGACTCTTGTTTTGGAGTTAGTTTGTGCTTATTTTGTGGAAATG
ATTGTTTTTCCTAGTAACAAAGCAG**Y**GCAGTTCACAAAGCAGTAAATGCTTC
AGCTCTCTTTTTCAGTTAACTATATTGAAATTAAATTCACTTTgatttttcttccctctcttg
aga
F 5'-3' = aaactttacagtcctttgggata
R 5'-3' = tctcaagagagggaagaaaaatc

M311 = EIFIA STS 39 (460 bp) **G to T** at position 304
(Group X)
CgagaacagcctaaccaacaTGGTGAAACCCCATCTCTGCTAAAAATATAAAAATTAGC
CAGGCATGGTAGTGCACACCTGTAGTCCCAGCTACTCAGGAGGCTGAGGCAG
GATAATCACTTGGACCCAGGAGACAGAGGTTGCAGTGAACCGAGATTGCACC
ACTGCACTCCAGCCTGGGCAATAGAGCGAGACTCCATCTCAAAAAAAAAAA
AAAAATTACAAAGGCTAAACTTTGGAAAGTCTAAGACAGACATAGGTGATGG
TCACACACTCCATTGAGAACCATTGTTCTACATCAGG**K**TTCTCTACAGCTTTT
GTTTTACCAACATGTTTATTAAGATTGTTTCCAGACTGTTCAGAGGAGTAGAA
GGATTTTTAAATTTATTTGTA**A**ACATTCAAATACTCACCAACAATATTGTACA
ATTTACAGTTTTTctctgcttcatctatcacaccc
F 5'-3' = cgagaacagcctaaccaaca
R 5'-3' = gggtgtgatagatgaagcagag

M312 = EIF1AY STS40a, **A to T** at position 49,
(Group VII)
gtttccagactgttcagaggagTAGAAGGATTTTTAAATTTATTTGTA**W**ACATTCAAATAC
TCACCAACAATATTGTACAATTTACAGTTTTTCTCTGCTTCATCTATCACACCC
ATCCTTCTATTCATCTGATATTACACCTTATATTTTGGCACATTTCCAAACTAT

TACTTACACTTTGAGTTGAAGAAAATAAACTGAGTCCTTAATTGTATTGTATA
TATGCATTTATAAATTTTTACAACATAAAGTACTCTATATTTACAAAATTTTTT
AGTTTTTTTTTT**T**CTTTGGAATTGTTTCTGAGTAGTACTTAGTAACACTACTCTA
ATGTAATATAAATTTTAAAGTATACCCAAAAAGAAAATGAAAAGAGATGAA
AAATGCATTGTTCTTGTGATCCCAGGAAATCTGAGAC**A**GGTCTCAGTTAATTT
acaaagttgattttgccaaagt
F 5'-3' = gtttccagactgttcagaggag
R 5'-3' = actttggcaaaatcaactttgt

M313 = EIFIA STS 40b **Homopolymer 9T's to 10T's** at position 288

gtttccagactgttcagaggagTAGAAGGATTTTTAAATTTATTTGTA**W**ACATTCAAATAC
TCACCAACAATATTGTACAATTTACAGTTTTTCTCTGCTTCATCTATCACACCC
ATCCTTCTATTCATCTGATATTACACCTTATATTTTGGCACATTTCCAAACTAT
TACTTACACTTTGAGTTGAAGAAAATAAACTGAGTCCTTAATTGTATTGTATA
TATGCATTTATAAATTTTTACAACATAAAGTACTCTATATTTACAAAATTTTTT
AGTTTTTTTTTT**T**CTTTGGAATTGTTTCTGAGTAGTACTTAGTAACACTACTCTA
ATGTAATATAAATTTTAAAGTATACCCAAAAAGAAAATGAAAAGAGATGAA
AAATGCATTGTTCTTGTGATCCCAGGAAATCTGAGAC**A**GGTCTCAGTTAATTT
acaaagttgattttgccaaagt
For 5'-3' = gtttccagactgttcagaggag
Rev 5'-3' = actttggcaaaatcaactttgt

M314 = EIFIA STS 40c (623 bp) **A to C** at position 419.
(Group VI)
GtttccagactgttcagaggAGTAGAAGGATTTTTAAATTTATTTGTA**A**ACATTCAAATA
CTCACCAACAATATTGTACAATTTACAGTTTTTCTCTGCTTCATCTATCACACC
CATCCTTCTATTCATCTGATATTACACCTTATATTTTGGCACATTTCCAAACTA
TTACTTACACTTTGAGTTGAAGAAAATAAACTGAGTCCTTAATTGTATTGTAT
ATATGCATTTATAAATTTTTACAACATAAAGTACTCTATATTTACAAAATTTTT
TAGTTTTTTTTTT**T**CTTTGGAATTGTTTCTGAGTAGTACTTAGTAACACTACTCT
AATGTAATATAAATTTTAAAGTATACCCAAAAAGAAAATGAAAAGAGATGA
AAAATGCATTGTTCTTGTGATCCCAGGAAATCTGAGAC**M**GGTCTCAGTTAAT
TTACAAAGTTGATTTTGCCAAAGTTGAGGACGCACCCATGACACAGCCTCGG
GAAGCCCTGAGGACATGTACCCAAGGTGTTTGGGGCACAGCTTGGTTTACTA
CATCTTCAGGGAGACATGAGACATCAATCAATATATGTGAAAAGAACGTTGG
TTCAGTTTGGAAAGGgagggcatcttgttagcctt
F 5'-3' = gtttccagactgttcagagg
R 5'-3' = aaggctaacaagatgccctc

M315 = EIFIA STS 41 (512 bp) **A to C** at position 395 STS also contains M314
GttcttgtgatcccaggaaatCTGAGAC**A**GGTCTCAGTTAATTTACAAAGTTGATTTTGCC
AAAGTTGAGGACGCACCCATGACACAGCCTCGGGAAGCCCTGAGGACATGT
ACCCAAGGTGTTTGGGGCACAGCTTGGTTTACTACATCTTCAGGGAGACATG
AGACATCAATCAATATATGTGAAAAGAACGTTGGTTCAGTTTGGAAAGGGAG

GGCATCTTGTTAGCCTTTCTAAAGGAGGCAGTCAGCTATGCATCTAACTCAAT
GAGCGAAAGGATAACTTTTGAATAGAATGGGAGGCCGGTTTGTCTTAAGCAG
TTTCCACCTTGAGTTTTTCATAGTAATTTTGGGGGCCAAAGATATTTTCGTTTC
ACATTCTAATATTTTCTTC**M**TGTACCTCCCTTTGGGGACCCTGAGCCAGAGGT
TTTTTGGGGGATTAAACAGAATTGGCATTTACTTCATGTTGCAATAACCAAAA
GCATAAATAttttgttgtagattaagggcaa
F 5'-3' = gttcttgtgatcccaggaaat
R 5'-3' = ttgcccttaatctacaacaaaa

M316 = EIFIA STS 42 (512 bp nominal) **5T's to 6T's** at position 201
Group V
AattggcatttacttcatgttgcAATAACCAAAAGCATAAATATTTTGTTGTAGATTAAGGgc
aaatctgaacatttccacAGTTGGTGGCCTTGGAGGCCTCTTTGGAAAATTCAGAGAACC
TATCCAGACTACCTAGTGGAACACAAAGCTACAAACACAGATGTTAGAATAA
GGATCTAGACATGGCTAAGATTTTT**T**CTCAGGGAGTGGGGGGGAGTATCTTA
GAGTTATGCCATTTCCTTTGGAACTAGGCCCATTAAGGTAACGGGAAGGAAT
GTAAAGACAATGGCTATTAAAGGAAGTTTAGTTTCTTTTGAGTTTCTTTTGCT
TATTACAAGAGAACACTGTAGATTTATAGATGTTCTAGTTTTACTTCTGTGAC
TACATGGACTCAGAATTTGGTTACGACCATATTTATCCCATTTTTAAAGGAAT
TACATCTATTTTGTCTGTGTCCACCCTCAGAATATAAGATCTGTAACCACTACc
acaaaaggaagtaaggacatg
F 5'-3' = aattggcatttacttcatgttgc
R 5'-3' = catgtccttacttcctttgtg

M317 = EIFIA STS 44 (523 bp nominal) **–2bp Deletion of GA** at position 400
(Group VIII)
TggttctacagttgggattttgGCCATCATCAACCAAGAAGAGAAATTCATTTAGTGTGTA
GTTTCTGAAAGCAAACTGATTTATTTTCATTGTTTTAAAGTATTTATTTCTTTA
AAAGCTGAGGACACTGAATTACCTTAAGTTAAATGTTAATACTTTATTGTTTT
GATGTAATGGAACTTAAGGATAAAAGACCATAATATTTGCTGTTAAAATAAA
TAAACGAGTGCCTTTCCTACTGTGATAACGTCAAGTAATTGGATATTTTGAAT
ACATTTCTGCCTGATAATCATGCTGGGTTCTAATAAGCCCTACTTCCACCTAA
TCTGTTTACAGTCTTTTGGTATGTTTCAGTTACTTAGATGGTCTCATAAGGTTT
CTGATACAATTTGAAGACA**GA**AATCTGCATTTAGAATCAGAAAACATGGAC
ATATTTTTCATATTTATCTAGTCATATGTAATTTTATGCTAACATTGATAGTTT
ATAAATCCTTTTCATCCTttgtgcctcggttattaagg
F 5'-3' = tggttctacagttgggattttg
R 5'-3' =ccttaataaccgaggcacaa

M318 = EIF1AY STS20d, **T to C**, at position 353 Group VI
Catggtccaagcaatttatttt**T**GTGAGTTCCCAAAATAATTTATACAGCAATGATTCATGT
GACAATGTGAATAAATAGAAAAAGTCTTTGATAACTTTTAGATTTACTTTTAA
AGAATAATTTGTTTGTTTAACTTCTGTTGTATTCCTACCAGAAATGTTTACTCT
GATATTAGTATTGAAGAAACCAGACAAATCTAATATATAACACAAATGGTCT
TGACTCAGATGTTAATGCTGTGAAAGAATGAAAAATCTGGGAATTACTTTAG
CTTAAAAGAGATTGATCGGTGCATA**T**CCCTT**C**GTTAGGTTTTGGATTGGGGGA

AATAGTTTTAGGTGGTACTAGGAAAA**Y**TGGAATATGGAATATGTTAGAAACT
CTATTTGTTAGTAATACCACATCAGGTAGTTT**T**ATAAATTACACTGATTAAAA
GTCTCTACTACTCAGATTTTTAATTAAAATAATAAAAACTTATTTTTGG**C**TGA
Gctctgtggaagtattagccagc
F 5'-3' = catggtccaagcaatttatttttg
Rev 5'-3' = gctggctaatacttccacagag

M319 = UTY1 exon 14b, **T to A** at position 124. Group VI
Gtaaaactcagatatatacatcccatg**A**AATATACACAGAAACTATAAATTAGCATTAATATC
CTCTAAAATGATACTGTAGTAAAGAAATATTCTCAAACTGTTGGTAAATTTTA
GAGAAAA**W**AAAAATATTATACATACTTGCTGCATTAAGACAAACTG**A**CTTTC
TAACTGTTCCAGCTGATGCTTCTGTGCTGGATTTAAATTATCTCTATTTGCTCG
CAGTTGTTCCAAGTGCTAGAAGAAAAGAGATTAATATAATCAAAGTTTAATC
TAAAATTTAAGACAATATAAGGCAACTCCTCACTAAAAAGACTACACAGAAC
CTTTGCAGGATGAAAGACAGTGATTCCTAATGAAcgttaagatagtgattctttttttt
F 5'-3' = gtaaaactcagatatatacatcccatg
Rev 5'-3': aaaaaaaaagaatcactatcttaacg
M320 = DBY STS08, (444 bp) T to G at position 60
Group VI
tgaggtggaatgtatcagtataccAATTAATATTTTTGAAAGAGCTCTTTTAGGTTAAT**K**TA
AGTACAGCAATTTCTCATGTAATGTTTAGGGAGTTTATTCTAACCTAGGCAAA
CGGCATGCTATCACAAGAAAGGTTTAAAGCTTTGATAAAATGGGGGAGATTT
AATCAGTTTTTTTAATGCCTGCTATAAAAATTTGAAATATTAGAATGGCCGAC
CATGGCAGTGACCAGGCCTCACTACAGGCCTGGTTGGATTCTGGTCTTTAATG
CATGCTAGTGTTGATGTTTTTTGGTCAAGAACGGTTTAAACAGGAAGGATTGT
GCAGCAGGCTTTAATTTAATGTAGATTCATACTGCTCTGTTAAAGCTGCATTG
AAATGTTAAAATGGCTTACACTTGCAGACTTTGCAAATCTTaagactaacaaatccttgaa
atca
For 5'-3'=tgaggtggaatgtatcagtatacc
Rev 5'-3'=tgatttcaaggatttgttagtctt

M321 = DBY STS08, (444 bp) C to T at position 171
group VI
tgaggtggaatgtatcagtataccAATTAATATTTTTGAAAGAGCTCTTTTAGGTTAATTAA
GTACAGCAATTTCTCATGTAATGTTTAGGGAGTTTATTCTAACCTAGGCAAAC
GGCATGCTATCACAAGAAAGGTTTAAAGCTTTGATAAAATGGGGGAGATTTA
AT**Y**AGTTTTTTTAATGCCTGCTATAAAAATTTGAAATATTAGAATGGCCGACC
ATGGCAGTGACCAGGCCTCACTACAGGCCTGGTTGGATTCTGGTCTTTAATGC
ATGCTAGTGTTGATGTTTTTTGGTCAAGAACGGTTTAAACAGGAAGGATTGTG
CAGCAGGCTTTAATTTAATGTAGATTCATACTGCTCTGTTAAAGCTGCATTGA
AATGTTAAAATGGCTTACACTTGCAGACTTTGCAAATCTTaagactaacaaatccttgaaat
ca
For 5'-3'=tgaggtggaatgtatcagtatacc
Rev 5'-3'=tgatttcaaggatttgttagtctt

Footnote:
STS sequences (one strand only) for polymorphic Y sequences.
Primer regions = lower case; Reverse compliment made to generate 5'-3' Reverse PCR primer sequence for complimentary strand.
IUB code defines polymorphic site
R = A or G (puRine)
Y = C or T (pYrimidine)
K = G or T (Keto)
M = A or C (aMino)
S = G or C (Strong-3H bonds)
W = A or T (Weak-2H bonds)
H = A, C or T

Markers M1, M29, M40, M46, M130, M167, M176, M177, M222, M236, M288 are unassigned in TABLE 1.